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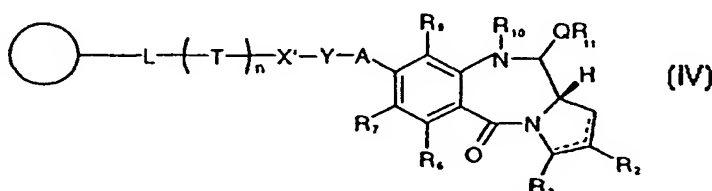
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(21) International Application Number: PCT/GB99/02836 (22) International Filing Date: 27 August 1999 (27.08.99) (30) Priority Data: 9818730.5 27 August 1998 (27.08.98) GB (71) Applicant (for all designated States except US): THE UNIVERSITY OF PORTSMOUTH HIGHER EDUCATION CORPORATION [GB/GB]; University House, Winston Churchill Avenue, Portsmouth PO1 2UP (GB). (72) Inventors; and (75) Inventors/Applicants (for US only): THURSTON, David, Edwin [GB/GB]; University of Portsmouth, St Michael's Building, White Swan Road, Portsmouth, Hampshire PO1 2UP (GB). HOWARD, Philip, Wilson [GB/GB]; University of Portsmouth, St Michael's Building, White Swan Road, Portsmouth, Hampshire PO1 2DT (GB). (74) Agents: WATSON, Robert, J. et al.; Mewburn Ellis, York House, 23 Kingsway, London WC2B 6HP (GB).		(81) Designated States: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZA, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SL, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG). Published Without international search report and to be republished upon receipt of that report.	

(54) Title: COLLECTIONS OF COMPOUNDS



(57) Abstract

A compound of formula (IV): O is a solid support; L is a linking group or a single bond; X' is selected from CO, NH, S, or O; A is O, S, NH, or a single bond; R₂ and R₃ are independently selected from: H, R, OH, OR, =O, =CH-R, =CH₂, CH₂-CO₂R, CH₂-CO₂H, CH₂-SO₂R, O-SO₂R, CO₂R, COR, CN and there is optionally a double bond between C1 and C2 or C2 and C3; R₆, R₇, and R₉ are independently selected from H, R, OH, OR, halo, nitro, amino, Me₃Sn; R₁₁ is either H or R; Q is S, O or NH; R₁₀ is a nitrogen protecting group; and Y is a divalent group such that HY = R, and other related compounds and collections of compounds.

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COLLECTIONS OF COMPOUNDS

This invention relates to collections of pyrrolobenzodiazepines, to methods of synthesizing these compounds on solid supports, and to compounds of utility therein. This invention further relates to methods for identifying and isolating pyrrolobenzodiazepine compounds with useful and diverse activities from such collections.

Background to the invention

Compounds having biological activity can be identified by screening collections of compounds (i.e. libraries of compounds) produced through synthetic chemical techniques. Such screening methods include methods wherein the library comprises a plurality of compounds synthesized at specific locations on the surface of a solid support where a receptor is appropriately labelled to identify binding to the compound, e.g., fluorescent or radioactive labels. Correlation of the labelled receptor bound to the support with its location on the support identifies the binding compound (US 5,143,854).

Central to these methods is the screening of a multiplicity of compounds in the library and the ability to identify the structures of the compounds which have a requisite biological activity. In order to facilitate synthesis and identification, the compounds in the library are typically formed on solid supports. Usually each such compound is covalently attached to the support via a cleavable or non-cleavable linking arm. The libraries of compounds can be screened either on the solid support or as cleaved products to identify compounds having good biological activity.

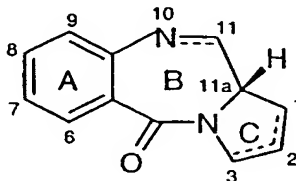
Background to the Invention

A large number of both synthetic and naturally occurring low molecular weight ligands are known that interact with DNA via a number of different mechanisms, including covalent or non-covalent interaction in the minor or major grooves, intercalation between base pairs or other types of non-

specific interactions.

A particular class of compounds which interacts with the minor groove are the pyrrolobenzodiazepines (PBDs). PBDs have the ability to recognise and bond to specific sequences of DNA; the most preferred sequence is PuGpu (Purine-Guanine-Purine). The first PBD antitumour antibiotic, anthramycin, was discovered in 1965 (Leimgruber et al., 1965 *J. Am. Chem. Soc.*, **87**, 5793-5795; Leimgruber et al., 1965 *J. Am. Chem. Soc.*, **87**, 5791-5793). Since then, a number of naturally occurring PBDs have been reported, and over 10 synthetic routes have been developed to a variety of analogues (Thurston et al., 1994 *Chem. Rev.* **1994**, 433-465). Family members include abbeymycin (Hochlowski et al., 1987 *J. Antibiotics*, **40**, 145-148), chicamycin (Konishi et al., 1984 *J. Antibiotics*, **37**, 200-206), DC-81 (Japanese Patent 58-180 487; Thurston et al., 1990, *Chem. Brit.*, **26**, 767-772; Bose et al., 1992 *Tetrahedron*, **48**, 751-758), mazethramycin (Kuminoto et al., 1980 *J. Antibiotics*, **33**, 665-667), neothramycins A and B (Takeuchi et al., 1976 *J. Antibiotics*, **29**, 93-96), porothramycin (Tsunakawa et al., 1988 *J. Antibiotics*, **41**, 1366-1373), prothracarcin (Shimizu et al., 1982 *J. Antibiotics*, **29**, 2492-2503; Langley and Thurston, 1987 *J. Org. Chem.*, **52**, 91-97), sibanomicin (DC-102) (Hara et al., 1988 *J. Antibiotics*, **41**, 702-704; Itoh et al., 1988 *J. Antibiotics*, **41**, 1281-1284), sibiromycin (Leber et al., 1988 *J. Am. Chem. Soc.*, **110**, 2992-2993) and tomamycin (Arima et al., 1972 *J. Antibiotics*, **25**, 437-444).

PBDs are of the general structure:

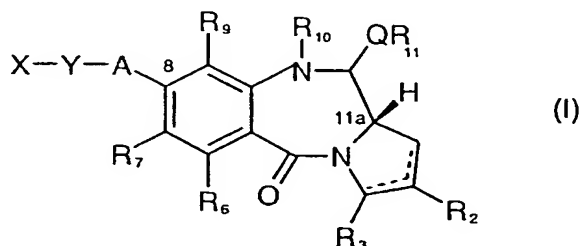


They differ in the number, type and position of substituents, in both their aromatic A rings and pyrrolo C rings, and in the

degree of saturation of the C ring. There is either an imine (N=C), carbinolamine (NH-CH(OH)) or a carbinolamine methyl ether (NH-CH(OMe)) at the N10-C11 position which is the electrophilic centre responsible for alkylating DNA. All of the known natural products have an (S)-configuration at the chiral C11a position which provides them with a right-handed twist when viewed from the C ring towards the A ring. This gives them the appropriate three-dimensional shape for isohelicity with the minor groove of B-form DNA, leading to a snug fit at the binding site (Kohn, 1975 In *Antibiotics III*. Springer-Verlag, New York, pp. 3-11; Hurley and Needham-VanDevanter, 1986 *Acc. Chem. Res.*, **19**, 230-237). Their ability to form an adduct in the minor groove enables them to interfere with DNA processing, hence their use as antitumour agents.

Disclosure of the Invention

A first aspect of the present invention relates to compounds of formula (I):



wherein:

X is selected from COOH, NHZ, SH, or OH, where Z is either H or an amine protecting group;

A is O, S, NH, or a single bond;

R₂ and R₃ are independently selected from: H, R, OH, OR, =O, =CH-R, =CH₂, CH₂-CO₂R, CH₂-CO₂H, CH₂-SO₂R, O-SO₂R, CO₂R, COR and CN, and there is optionally a double bond between C₁ and C₂ or C₂ and C₃;

R₆, R₇, and R₉ are independently selected from H, R, OH, OR, halo, nitro, amino, Me₃Sn;

R₁₁ is either H or R;

Q is S, O or NH;

R₁₀ is a nitrogen protecting group;

where R is a lower alkyl group having 1 to 10 carbon atoms, or an aralkyl group (i.e. an alkyl group with one or more aryl substituents), preferably of up to 12 carbon atoms, whereof
5 the alkyl group optionally contains one or more carbon-carbon double or triple bonds, which may form part of a conjugated system, or an aryl group, preferably of up to 12 carbon atoms; and is optionally substituted by one or more halo, hydroxy, amino, or nitro groups, and optionally contains one or more
10 hetero atoms, which may form part of, or be, a functional group; and

Y is a divalent group such that HY = R.

These compounds are useful in the synthesis of collections of
15 pyrrolobenzodiazepines. Compounds of formula I can be attached to a solid support, e.g. via a connecting link which may comprise a chain of combinatorial units. Without N10-protection, the risk of undesirable side reactions with the imine bond during the coupling step would be greater.

20 If R is an aryl group, and contains a hetero atom, then R is a heterocyclic group. If R is an alkyl chain, and contains a hetero atom, the hetero atom may be located anywhere in the alkyl chain, e.g. -O-C₂H₅, -CH₂-S-CH₃, or may form part of, or
25 be, a functional group, e.g. carbonyl, hydroxy.

R and HY groups are preferably independently selected from a lower alkyl group having 1 to 10 carbon atoms, or an aralkyl group, preferably of up to 12 carbon atoms, or an aryl group,
30 preferably of up to 12 carbon atoms, optionally substituted by one or more halo, hydroxy, amino, or nitro groups. It is more preferred that R and HY groups are independently selected from lower alkyl groups having 1 to 10 carbon atoms optionally substituted by one or more halo, hydroxy, amino, or nitro
35 groups. It is particularly preferred that R or HY are unsubstituted straight or branched chain alkyl groups, having 1 to 10, preferably 1 to 6, and more preferably 1 to 4, carbon atoms, e.g. methyl, ethyl, propyl, butyl.

Alternatively, R_6 , R_7 , and R_8 may preferably be independently selected from R groups with the following structural characteristics:

- (i) an optionally substituted phenyl group;
- 5 (ii) an optionally substituted ethenyl group;
- (iii) an ethenyl group conjugated to an electron sink.

The term 'electron sink' means a moiety covalently attached to a compound which is capable of reducing electron density in
10 other parts of the compound. Examples of electron sinks include cyano, carbonyl and ester groups.

The term 'nitrogen protecting group' (or 'amine protecting group') has the meaning usual in synthetic chemistry,
15 particularly synthetic peptide chemistry. It means any group which may be covalently bound to the nitrogen atom of the pyrrolobenzodiazepine (or amine) grouping, and permits reactions to be carried out upon the molecule containing this grouping without its removal. Nevertheless, it is able to be
20 removed from the nitrogen atom without affecting the remainder of the molecule. Suitable nitrogen protecting groups for the present invention include Fmoc (9-fluorenylmethoxycarbonyl), Nvoc (6-nitroveratryloxycarbonyl), Teoc (2-trimethylsilylethyloxycarbonyl), Troc (2,2,2-
25 trichloroethyloxycarbonyl), Boc (t-butyloxycarbonyl), CBZ (benzyloxycarbonyl), Alloc (allyloxycarbonyl), and Psec (2(-phenylsulphonyl)ethyloxycarbonyl). Other suitable groups are described in Protective Groups in Organic Synthesis, T Green and P Wuts, published by Wiley, 1991, which is incorporated
30 herein by reference. It is preferred that the nitrogen protecting group has a carbamate functionality where it binds to the nitrogen atom at the 10 position of the PBD ring structure.

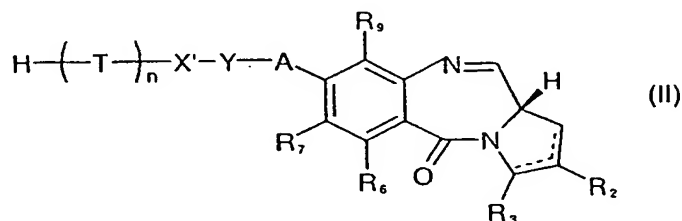
35 R_9 is preferably an electron donating group. 'Electron donating group' means a moiety covalently attached to a compound which is capable of increasing electron density in other parts of the molecule. Examples of electron donating

groups useful in the present invention include alkyl, amine, hydroxyl, alkoxy and the like.

In compounds of formula I, Q is preferably O, and R₁₁ is preferably H. Independently, R₆ and R₇ are preferably H, and R₇ is preferably an alkoxy group, and more preferably methoxy or ethoxy. It is further preferred that if there is a double bond in the C ring, it is between C2 and C3. In this case, R₂ and R₃ are preferably H.

-Y-A- is preferably an alkoxy chain, preferably an ethoxy chain.

A second aspect of the present invention relates to compounds of formula II :



wherein Y, A, R₇, R₂, R₃, R₆, and R₉ are as defined in the first aspect of the invention;

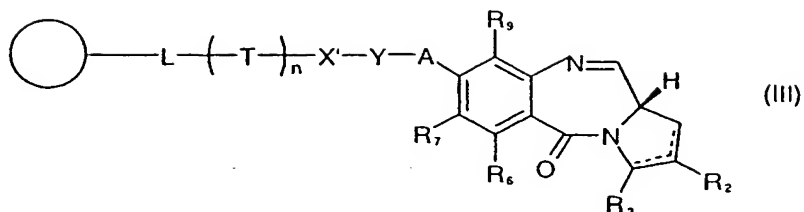
X' is CO, NH, S or O;

T is a combinatorial unit;

and n is a positive integer, where if n is greater than 1, each T may be different.

It is preferred that X' is either CO or NH. n may preferably be from 1 to 16, and more preferably from 3 to 14.

A third aspect of the present invention relates to compounds of the formula III:



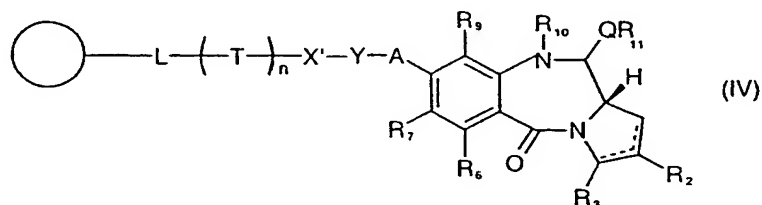
wherein X' , Y , A , R_7 , R_2 , R_3 , R_6 , R_9 , and T are as defined in the second aspect of the invention;

n is zero or a positive integer;

L is a linking group, or less preferably a single bond;

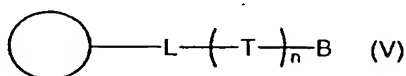
5 and O is a solid support, where if n is greater than 1, each T may be different.

A fourth aspect of the present invention relates to compounds of the formula IV:



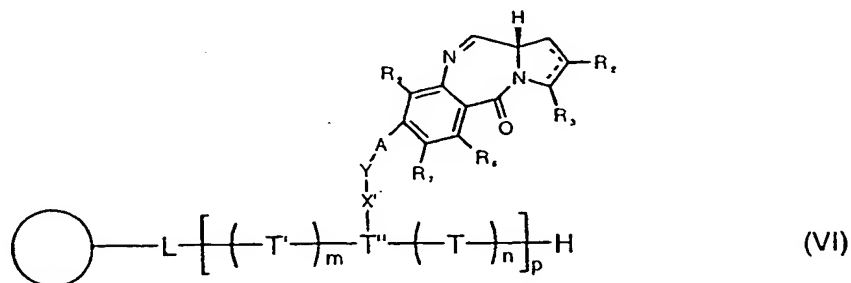
10 wherein X' , Y , A , R_7 , R_2 , R_3 , R_6 , R_9 , T , n , L and O are as defined in the third aspect of the invention, and R_{10} , R_{11} , and Q are as defined in the first aspect of the invention.

15 A fifth aspect of the present invention relates to a method of making compounds of formula IV as described in the fourth aspect of the invention by reacting compounds of formula I with compounds of formula V:



wherein O, L, T and n are as defined in the fourth aspect of the invention, and B is H or an atom or group for providing a functional group capable of reaction with X.

- 5 A sixth aspect of the present invention relates to compounds of formula VI:



wherein O, L, X', Y, A, R₂, R₃, R₆, R₇, R₉ and T are as defined in the second aspect of the invention;

- 10 n and m are positive integers, or one of them may be zero;

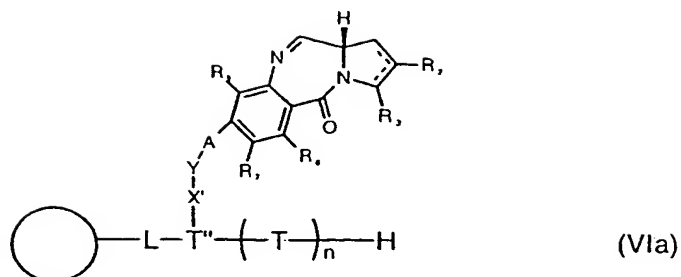
T' is a combinatorial unit, where each T' may be different if m is greater than 1;

T'' is a combinatorial unit which provides a site for the attachment of X'; and

- 15 p is a positive integer, where if p is greater than 1, for each repeating unit, the meaning of X', Y, A, R₂, R₃, R₆, R₇, R₉, T, T', T'' and the values of n and m are independently selected.

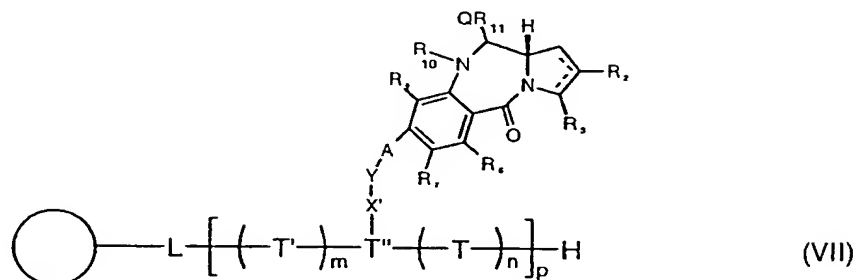
- 20 For example, if X' is CO then the site on T'' may be NH, and if X' is NH, S or O, then the site on T'' may be CO.

In a preferred aspect of the sixth aspect of the present invention, the compound is of formula (VIa):



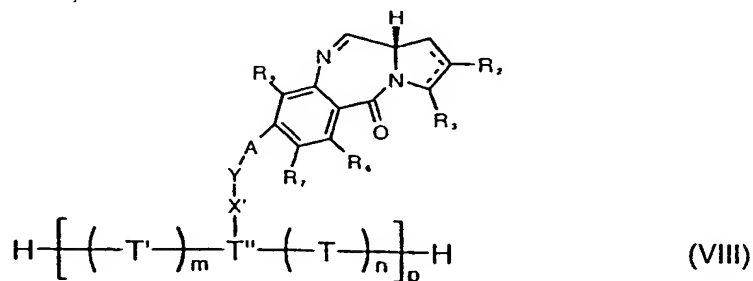
wherein O, L, X', Y, A, R₂, R₃, R₆, R₇, R₉, T, T'' are as defined above.

A seventh aspect of the present invention relates to compounds of formula **VII**:



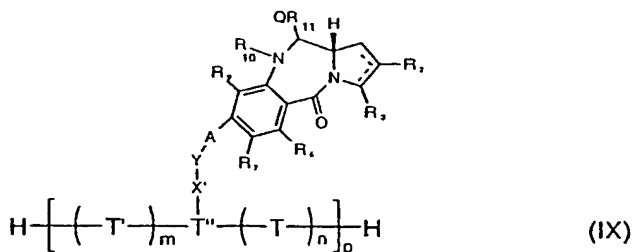
wherein O, L, X', Y, A, R₂, R₃, R₆, R₇, R₉, T, T', T'', n, m and p are as defined in the sixth aspect and Q, R₁₀, and R₁₁, are as defined in the first aspect of the invention, where if p is greater than 1, for each repeating unit the meanings of X', Y, A, R₂, R₃, R₆, R₇, R₉, T, T', T'', Q, R₁₀, R₁₁ and the values of n and m are independently selected.

An eighth aspect of the present invention relates to compounds of formula **VIII**:



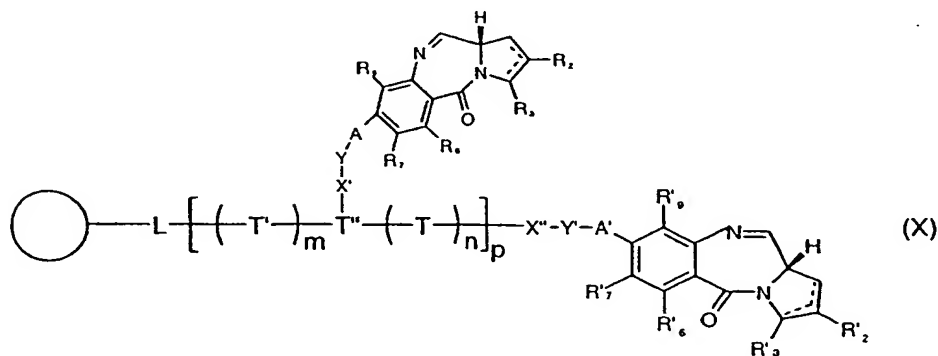
wherein X' , Y , A , R_2 , R_3 , R_6 , R_7 , R_9 , T , T' , T'' , n , m and p are as defined in the sixth aspect of the invention, where if p is greater than 1, for each repeating unit the meanings of X' , Y , A , R_2 , R_3 , R_6 , R_7 , R_9 , T , T' , T'' and values of n and m are independently selected.

A ninth aspect of the present invention relates to compounds of formula IX:



wherein X' , Y , A , R_2 , R_3 , R_6 , R_7 , R_9 , Q , R_{10} , R_{11} , T , T' , T'' , n , m and p are as defined in the seventh aspect of the invention, where if p is greater than 1, for each repeating unit the meanings of X' , Y , A , R_2 , R_3 , R_6 , R_7 , R_9 , T , T' , T'' , Q , R_{10} , R_{11} and values of n and m are independently selected.

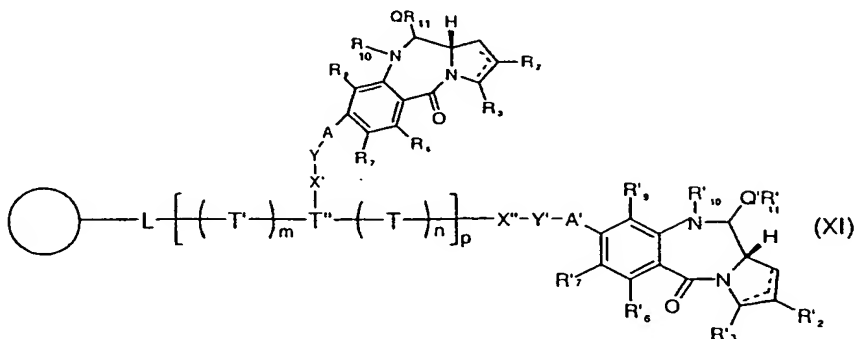
A tenth aspect of the present invention relates to compounds of formula X:



wherein O , L , X' , Y , A , R_2 , R_3 , R_6 , R_7 , R_9 , T , T' , T'' , n , m and p are as defined in the sixth aspect of the invention, and X'' , Y' , A' , R'_2 , R'_3 , R'_6 , R'_7 , and R'_9 are selected from the same possibilities as X' , Y , A , R_2 , R_3 , R_6 , R_7 , and R_9 , respectively, and where if p is greater than 1, for each repeating unit the

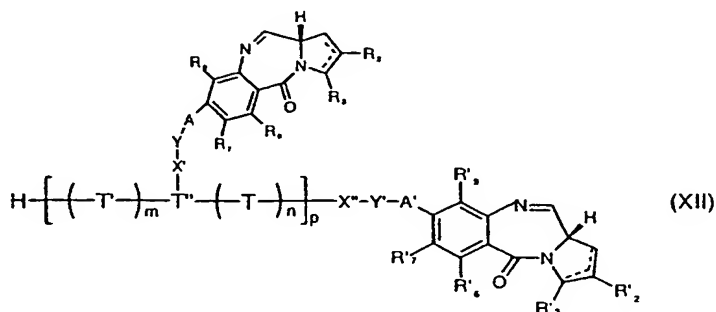
meaning of X' , Y , A , R_2 , R_3 , R_6 , R_7 , R_9 , T , T' , T'' and the values of n and m may be independently selected.

An eleventh aspect of the present invention relates to compounds of formula **XI**:



wherein O , L , X' , Y , A , R_2 , R_3 , R_6 , R_7 , R_9 , X'' , Y' , A' , R'_2 , R'_3 , R'_6 , R'_7 , R'_9 , T , T' , T'' , n , m and p are as defined in the tenth aspect of the invention, Q , R_{10} , and R_{11} , are as defined in the first aspect of the invention, and Q' , R'_{10} , R'_{11} , have the same definitions as Q , R_{10} , R_{11} , respectively, and where if p is greater than 1, for each repeating unit the meanings of X' , Y , A , R_2 , R_3 , R_6 , R_7 , R_9 , T , T' , T'' , Q , R_{10} , R_{11} and the values of n and m are independently selected.

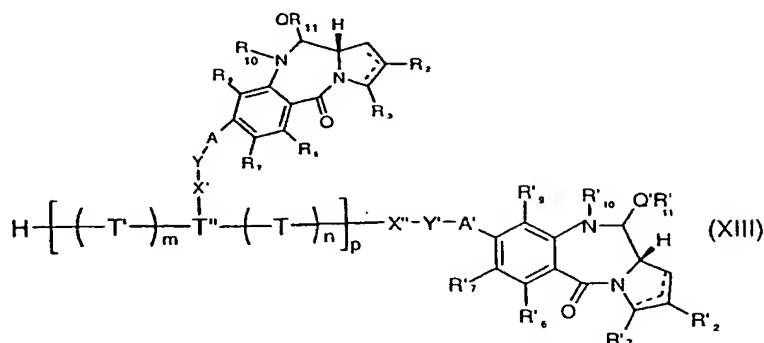
A twelfth aspect of the present invention relates to compounds of formula **XII**:



wherein X' , Y , A , R_2 , R_3 , R_6 , R_7 , R_9 , X'' , Y' , A' , R'_2 , R'_3 , R'_6 , R'_7 , R'_9 , T , T' , T'' , n , m and p are as defined in the tenth aspect of the invention, and where if p is greater than 1, for each repeating unit the meanings of X' , Y , A , R_2 , R_3 , R_6 , R_7 , R_9 ,

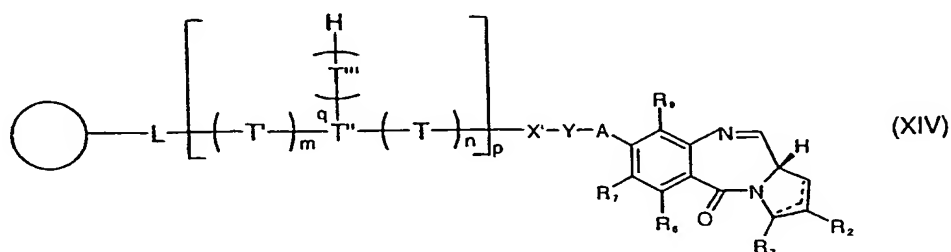
T, T', and T'' and the values of n and m may be independently selected.

A thirteenth aspect of the present invention relates to compounds of formula XIII:



wherein X', Y, A, R₂, R₃, R₆, R₇, R₉, Q, R₁₀, R₁₁, X'', Y', A', R'₂, R'₃, R'₆, R'₇, R'₉, Q', R'₁₀, R'₁₁, T, T', T'', n, m and p are as defined in the eleventh aspect of the invention, and where if p is greater than 1, for each repeating unit the meanings of X', Y, A, R₂, R₃, R₆, R₇, R₉, T, T', T'', Q, R₁₀, R₁₁ and the values of n and m may be independently selected.

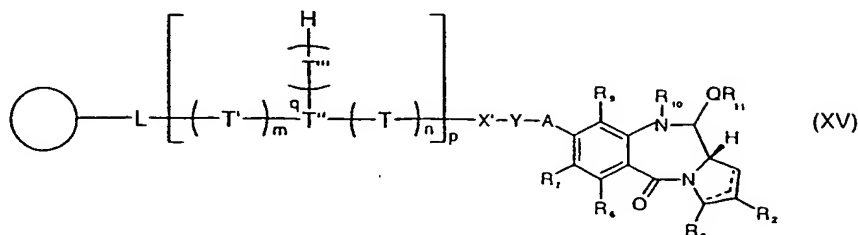
A fourteenth aspect of the present invention relates to compounds of formula XIV:



wherein O, L, X', Y, A, R₂, R₃, R₆, R₇, R₉, T, T', T'', n, m and p are as defined in the sixth aspect of the invention, and T''' and q are selected from the same possibilities as T and n respectively, and where if p is greater than 1, for each repeating unit the meaning of T, T', T'', T''' and the values of

n, m and q may be independently selected.

A fifteenth aspect of the present invention relates to compounds of formula **XV**:

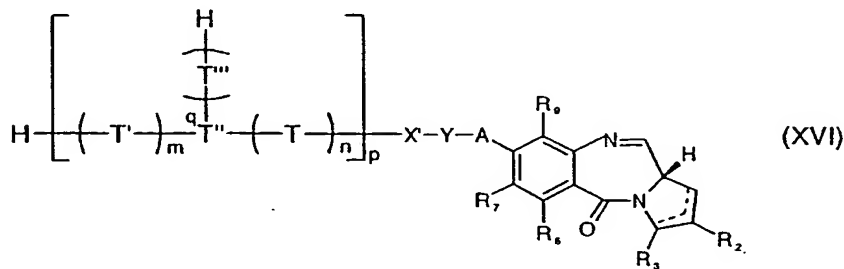


5

wherein O, L, X', Y, A, R₂, R₃, R₆, R₇, R₉, T, T', T'', T''', n, m, p and q are as defined in the fourteenth aspect of the invention, Q, R₁₀, and R₁₁, are as defined in the first aspect of the invention, and where if p is greater than 1, for each repeating unit the meanings of T, T', T'', T''' and the values of n, m and q may be independently selected.

10

A sixteenth aspect of the present invention relates to compounds of formula **XVI**:



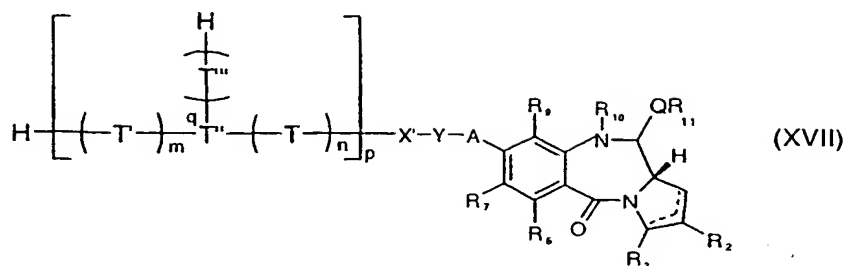
15

wherein X', Y, A, R₇, R₂, R₃, R₆, R₉, T, T', T'', T''', n, m, p and q are as defined in the fourteenth aspect of the invention, and where if p is greater than 1, the meanings of T, T', T'', T''' and values of n, m and q may be independently selected.

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A seventeenth aspect of the present invention relates to

compounds of formula XVII:



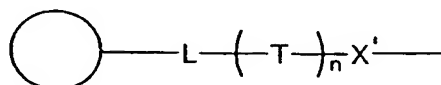
wherein X', Y, A, R₂, R₃, R₆, R₇, R₉, Q, R₁₀, R₁₁, T, T', T'', T''',
 n, m, p and q are as defined in the fourteenth aspect of the
 invention, and where if p is greater than 1, for each
 5 repeating unit the meanings of T, T', T'', T''' and the values
 of n, m and q may be independently selected.

Solid support

The term 'solid support' refers to a material having a rigid
 10 or semi-rigid surface which contains or can be derivatized to
 contain reactive functionalities which can serve for
 covalently linking a compound to the surface thereof. Such
 materials are well known in the art and include, by way of
 example, silicon dioxide supports containing reactive Si-OH
 15 groups, polyacrylamide supports, polystyrene supports,
 polyethyleneglycol supports, and the like. Such supports will
 preferably take the form of small beads, pins/crowns, laminar
 surfaces, pellets, disks. Other conventional forms may also
 be used.

Linker group

One class of linking groups suitable for the present
 application is one which provide in the structure:



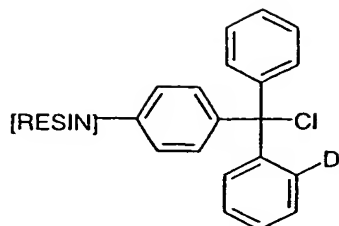
at least one covalent bond which can be readily broken by
 25 specific chemical reactions (or by light or changes in pH)

thereby providing for liberation of compounds free from the solid support. The methods employed to break the covalent bond are selected so as to be specific for the desired bond breakage thereby preventing unintended reactions from occurring elsewhere on the complex. The linking group is selected relative to the synthesis of the compounds to be formed on the solid support so as to prevent premature cleavage of this compound from the solid support as well as to limit interference by any of the procedures employed during compound synthesis on the support.

Examples of resins incorporating linking groups are set out in the table below, which also indicates the groups that can be immobilised thereon, along with the suggested cleavage methods for the linking group. Such resins are commercially available (e.g. from NovaBiochem). The table also indicates which of the linker groups are suitable for compounds where the N10 position is not protected, i.e. where the cleavage methods would not affect an N10-C11 imine bond.

Linker/Resin Type	Immobilises	Cleavage Method	Compatible with Non-Protected PBD
2-Chlorotrityl chloride	RNH ₂ , RCO ₂ H, ROH, RSH	1-50% TFA	Possibly
Trityl chloride	RNH ₂ , RCO ₂ H, ROH, RSH	1-5% TFA	Yes
2-Methoxytrityl chloride	RNH ₂ , RCO ₂ H, ROH, RSH	1-5%	Yes
Rink amide resin	RCO ₂ H	95% TFA	Yes
Sieber amide resin	RCO ₂ H	1% TFA	Yes
4-Sulfamyl-benzoyl	RCO ₂ H	Alkylation /amines	YES
Wang resin	ROH, AROH, RNH ₂ , RCO ₂ H	15-95% TFA or DDQ or CAN	Possibly
HMPB-BHA	ROH, AROH, RCO ₂ H	1% TFA	Possibly
Bromoethyl photolinker	RNH ₂ , RCO ₂ H, ROH, RSH	hν	YES
Hydroxy ethyl photolinker	RCO ₂ H	hν	YES
Aminoethyl photolinker	RCO ₂ H	hν	YES

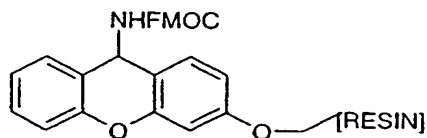
Structures



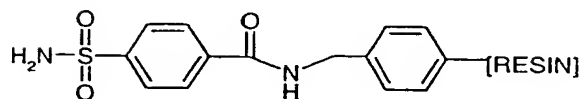
D = Cl: 2-chlorotrityl chloride type
 D = H: trityl chloride type
 D = OMe: 2-methoxytrityl chloride



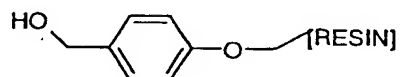
Rink amide type



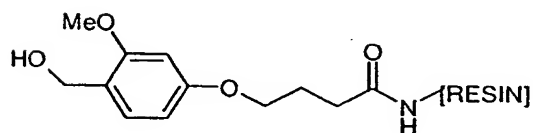
Sieber amide type



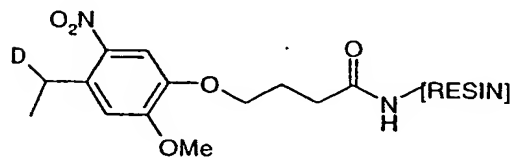
4-sulfamyl-benzoyl type



Wang type



HMPB-BHA type



D = NH₂: amino-ethyl type
 D = OH: hydroxy-ethyl type
 D = Br: bromo ethyl type

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For protected PBDs the most preferred linking group is the Rink linker, which is cleavable by TFA. The N-protected PBDs can then be deprotected using photolysis. For unprotected

PBDs the linking groups of choice are those which are photolabile.

It is also possible that the linking group is a simple functionality provided on the solid support, e.g. amine, and in this case the linking group may not be readily cleavable. This type of linking group is useful in the synthesis of large split and mix libraries which will be subjected to on-bead screening (see below), where cleavage is unnecessary. Such resins are commercially available from a large number of companies including NovaBiochem, Advanced ChemTech and Rapp Polymere. These resins include amino-Tentagel, and amino methylated polystyrene resin.

Combinatorial Unit

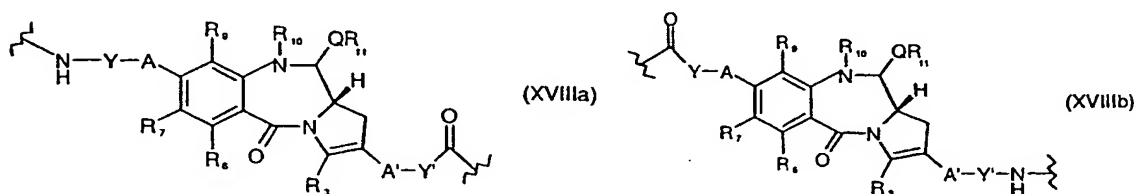
The term 'combinatorial unit' means any monomer unit which can be used to build a chain attached to the solid support, usually by a linking group. Examples of molecules suitable for such chain building are found in Schreiber et al. (*JACS*, 120, 1998, pp.23-29), which is incorporated herein by reference. An important example of a unit is an amino acid residue. Chains may be synthesised by means of amine-protected amino acids. Fmoc protected amino-acids are available from a number of sources, such as Sigma and Nova Biochem. Both natural and unnatural amino acids can be used, e.g. D- and L-amino acids and heterocyclic amino acids. In particular, heterocyclic amino acids of the type found in the construction of netropsin and distamycin are of interest because of their DNA-recognition properties.

Amine units can be used to make up peptoids: see Soth, M.J. and Nowick, J.S. 1997, Unnatural oligomer libraries, *Curr. Opin. Chem. Biol.* 1, no. 1: 120-129; Zuckermann et al., 1994, Discovery of Nanomolecular Ligands for 7-Transmembrane G-Protein-Coupled Receptors from a Diverse N-(Substituted)glycine Peptoid Library, *Journal of Medicinal Chemistry* 37: 2678-85; Figliozzi, GMR et al., 1996, Synthesis of N-substituted Glycine Peptoid Libraries, *Methods in*

Enzymology, 267: 437-47; Simon, R J et al., 1992, Peptoids: A Modular Approach to Drug Discovery, *Proc. Natl. Acad. Sci. USA*, 89:9367-71; which are all incorporated herein by reference.

Other combinatorial units include PNAs (peptidonucleic acids): P E Nielsen, et al, *Science*, 1991, 254, 1497; M Egholm, et al, *Nature*, 1993, 365, 566; M Egholm et al, *JACS*, 1992, 114, 1895; S C Brown, et al, *Science*, 1994, 265, 777; 5. K Saha, et al, *JOC*, 1993, 58, 7827; oligoureas: Burgess K, et al, 1995, Solid Phase Synthesis of Unnatural Biopolymers Containing Repeating Urea Units. *Agnew. Chem. Int. Examining Division. Engl* 34, no. 8:907; Burgess K, et al, 1997, Solid Phase Synthesis of Oligoureas; *Journal of the American Chemical Society* 119: 1556-64; and oligocarbamates: Moran E J et al, 1995, Novel Biopolymers for Drug Discovery. *Biopolymers (Peptide Science)*; John Wiley and Sons 37: 213-19; Cho C Y et al, 1993, An Unnatural Biopolymer. *Science* 261: 1303-5; Paikoff S F et al, 1996, The Solid Phase Synthesis of N-Alkylcarbamate Oligomers. *Tetrahedron Letters* 37, no. 32: 5653-56. All of these documents are incorporated herein by reference.

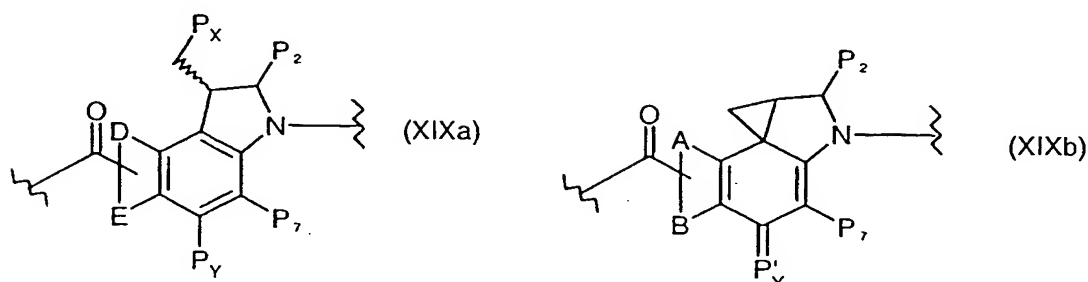
A type of combinatorial unit of particular relevance to the present invention is one based on the pyrrolobenzodiazepine structures; these are of general formulae **XVIIIa** and **XVIIIb**:



wherein R₃, R₆, R₇, R₈, R₉, R₁₀, R₁₁, Q, A and Y are as defined in the first aspect of the invention, and A' and Y' are independently selected from the possible groups for A and Y respectively.

A further type of particularly relevant combinatorial unit is one based on a cyclopropyl indole ("a CPI unit"). Such units

are known to interact covalently with the minor groove of DNA, being specific for AT. These units are of general formulae XIXa and XIXb:



wherein P_x (if present) is an electrophilic leaving group;

P_y (if present) is selected from NH-Prot, O-Prot, S-Prot, NO₂, NHOH, N₃, NHR, NRR, N=NR, N(O)RR, NHSO₂R, N=NPhR, SR or SSR, where Prot represents a protecting group;

P'_y (if present) is selected from NH, O and S;

D and E collectively represent a fused benzene or pyrrole ring (in either orientation), which is optionally substituted by up to respectively 3 or 1 additional groups independently selected from R, OH, OR, halo, nitro, amino, Me₃Sn, CO₂H, CO₂R; P₂ and P_z are independently selected from H, R, OH, OR, halo, nitro, amino, Me₃Sn.

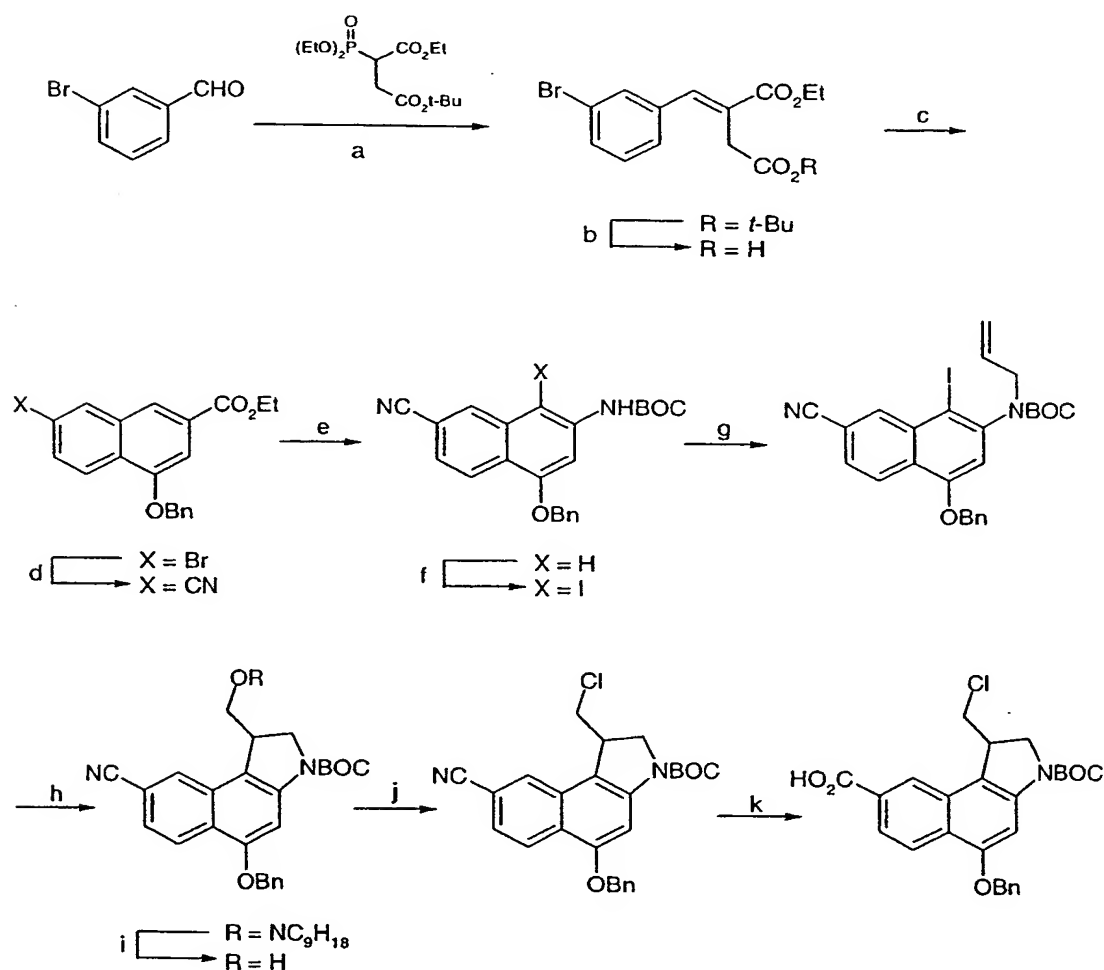
The preferences for R are as above. P_y is preferably NH-Prot, O-Prot, S-Prot and P_x is preferably halogen or OSO₂R. It is further preferred that the -CO₂- substituent is in the 2 or 3 position of the benzene ring or the 2 position of the pyrrole ring. P₂ and P_z are preferably H.

These compounds may be synthesised using the techniques described by Boger et al, *Chem. Rev.* **1997**, *97*, 787-828; Cava et al, Drost, K.J.; Cava, M.P. *J. Org. Chem.* **1991**, *56*, 2240-2244; Rawal, V.H.; Jones, R.J.; Cava, M.P. *J. Org. Chem.* **1987**, *52*, 19-28; and Aristoff *J. Med. Chem.* **1993**, *1992*, *57*, 6234-6239.

The following synthesis is provided as an example of the application of the Boger method.

Synthesis of a CPI Combinational Unit

The synthesis starts with a Wadsworth-Horner-Emmons condensation of 3-bromo-benzaldehyde with the Sargent phosphonate which predominantly provides the E-isomer, which in turn undergoes acid-catalyzed deprotection and Friedel-Crafts acylation. This generates the functionalised precursor, which is followed by 5-exo-trig aryl radical-alkene cyclization.



Reagents and conditions: a: NaH, Sargent phosphate; b; TFA; c: 1) Ac_2O -KOAc; 2) K_2CO_3 ; 3) BnBr, K_2CO_3 ; d: CuCN; e: 1) LiOH; 2) DPPA, $t\text{-BuOH}$; f: NIS; g: allyl Br, NaH; h: Bu_3SnH , TEMPO; i: Zn-HOAc; j: $\text{Ph}_3\text{P-CCl}_4$; k: NaOH.

Aromatic nucleophilic substitution, ester hydrolysis and Curtius rearrangements effected by treatment with DPPA are

followed by regioselective C4 iodination and N-alkylation with allyl bromide. The aryl radical-alkene cyclization by means of TEMPO as radical trap, as described in Boger synthesis of CBI, provides the tricyclic system that, after conversion to the primary chloride and base-catalyzed hydrolysis of the cyano group, gives the desired combinatorial unit.

The present invention relates to libraries, or collections, of compounds all of which are represented by a single one of the formulae I to IV, and VI to XVII. The diversity of the compounds in a library may reflect the presence of compounds differing in the identities of one or more of the substituent groups and/or in the identities of the combinatorial units T (when present). The number of members in the library depends on the number of variants, and the number of possibilities for each variant. For example, if it is the combinatorial units which are varied, and there are 3 combinatorial units, with 3 possibilities for each unit the library will have 27 compounds. 4 combinatorial units and 5 possibilities for each unit gives a library of 625 compounds. If for instance there is a chain of 5 combinatorial units with 17 possibilities for each unit, the total number of members in the library would be 1.4 million. A library may therefore comprise more than 1 000, 5 000, 10 000, 100 000 or a million compounds, which may be arranged as described below.

In the case of free compounds (formulae I, II, VIII, IX, XII, XIII, XVI, and XVII) the individual compounds are preferably in discrete volumes of solvents, e.g. in tubes or wells. In the case of bound compounds (formulae III, IV, VI, VII, X, XI, XIV and XV) the individual compounds are preferably bound at discrete locations, e.g. on respective pins/crowns or beads. The library of compounds may be provided on a plate which is of a suitable size for the library, or may be on a number of plates of a standard size, e.g. 96 well plates. If the number of members of the library is large, it is preferable that each well on a plate contains a number of related compounds from the library, e.g. from 10 to 100. One possibility for this type of grouping of compounds is where only a subset of the

combinatorial units, or substituents, are known and the remainder are randomised; this arrangement is useful in iterative screening processes (see below). The library may be presented in other forms that are well-known.

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A further aspect of the present invention is a method of preparing a diverse collection, or library of compounds as discussed above. If the diversity of the library is in the combinatorial units, then the library may be synthesised by the stepwise addition of protected combinatorial units to a PBD core, each step being interposed by a deprotection step. Such a method is exemplified later. Libraries of this type can be prepared by the method known as "split and mix" which is described in Furka, A; Sebestyen, F; Asgedom, M and Dibo, G; General Method of Rapid Synthesis of Multicomponent Peptide Mixtures; International Journal of Peptide and Protein Research; 1991, 37, 487-193, which is incorporated herein by reference. If the diversity of the library is in the substituent groups, the library may be synthesised by carrying out the same synthetic methods on a variety of starting materials or key intermediates, which already possess the necessary substituent patterns.

The present invention also relates to a method of screening the compounds of formula II, III, VI, VIII, X, XII, XIV and XVI to discover biologically active compounds. The screening can be to assess the binding interaction with nucleic acids, e.g. DNA or RNA, or proteins, or to assess the affect of the compounds against protein-protein or nucleic acid-protein interactions, e.g. transcription factor DP-1 with E2F-1, or estrogen response element (ERE) with human estrogen receptor (a 66 kD protein which functions as hormone-activated transcription factor, the sequence of which is published in the art and is generally available). The screening can be carried out by bringing the target macromolecules into contact with individual compounds or the arrays or libraries described above, and selecting those compounds, or wells with mixtures of compounds, which show the strongest effect.

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This effect may simply be the cytotoxicity of the compounds in question against cells or the binding of the compounds to nucleic acids. In the case of protein-protein or nucleic acid-protein interaction, the effect may be the disruption of the interaction studied.

The binding of the compounds to nucleic acids may be assessed by labelling oligomers which contain a target sequence, and measuring the amount of labelled oligomers that bind to the compounds tested. The labelling may either be radio-labelling, or alternatively be labels detectable under visible or ultra-violet light. If this latter form of screening is carried out on compounds bound to solid supports which are in separate locations, the screening for results can be carried out visually under a microscope. A similar technique is described in detail in "DNA-Binding ligands from peptide libraries containing unnatural amino acids", Lescrinier et al., *Chem Eur J*, 1998, 425-433. These techniques are particularly suited to a one-step screening of a complete library of compounds, especially a large library made by the "split and mix" method described above.

Protein-protein interactions can be measured in a number of ways, e.g. FRET (fluorescence resonance energy transfer) which involves labelling one of the proteins with a fluorescent donor moiety and the other with an acceptor which is capable of absorbing the emission from the donor; the fluorescence signal of the donor will be altered depending on the interaction between the two proteins. Another method of measuring protein-protein interactions is by enzymatic labelling, using, for example, horseradish peroxidase.

The screening process may undergo several iterations by selecting the most active compounds, or group of compounds, tested in each iteration; this is particular useful when testing arrays of wells which include mixtures of related compounds. Furthermore, if the wells contain compounds for which only a subset of the combinatorial units, or

substituents, are known, but the rest are randomised, subsequent iterations can be carried out by synthesising compounds possessing the selected known (and successful) combinatorial unit, or substituent, pattern, but with further specified combinatorial units, or substituents, replacing the previously randomised combinatorial units, or substituents, adjacent the already known pattern; the remaining combinatorial units, or substituents, are randomised as in the previous iteration. This iterative method enables the identification of active members of large libraries without the need to isolate every member of the library.

A further feature of this aspect is formulation of selected compound or compounds with pharmaceutically acceptable carriers or diluents.

In yet further aspects, the invention provides a pharmaceutical composition comprising a compound of formula **II**, **VIII**, **XII** or **XVI** and a pharmaceutically acceptable carrier or diluent; and the use of a compound of formula **II**, **VIII**, **XII** or **XVI** in the manufacture of a medicament for the treatment of a gene-based disease, or a bacterial, parasitic or viral infection. Gene-based disease include neoplastic disease, and Alzheimer's disease, and also include any disease susceptible to regulation of gene-expression.

Compounds of formula **II**, **VIII**, **XII** or **XVI** may be used in a method of therapy against a gene-based disease, such as cancer or Alzheimer's disease, or a viral, parasitic or bacterial infection.

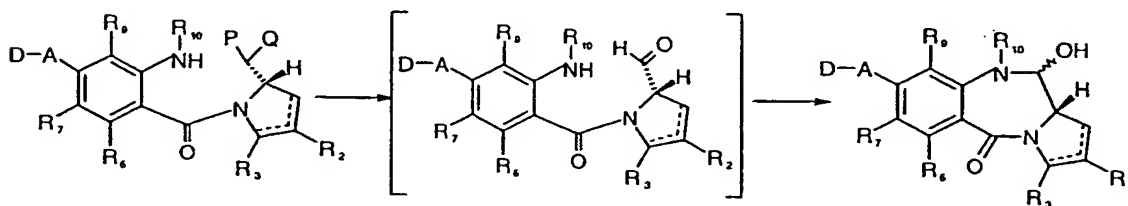
Another aspect of the present invention relates to the use of compounds of formula **III**, **VI**, **X** or **XIV** in diagnostic methods. A compound of formula **III**, **VI**, **X**, **XIV** which binds to an identified sequence of DNA or a protein known to be an indicator of a medical condition can be used in a method of diagnosis. The method may involve passing a sample, e.g. of appropriately treated blood or tissue extract, over an

immobilised compound of formula **III**, **VI**, **X**, **XIV**, for example in a column, and subsequently determining whether any binding of target DNA to the compound of formula **III**, **VI** or **X** has taken place. Such a determination could be carried out by passing a known amount of labelled target DNA known to bind to compound **III**, **VI** or **X** through the column, and calculating the amount of compound **III**, **VI**, **X** or **XIV** that has remained unbound.

A further aspect of the present invention relates to the use of compounds of formula **II**, **VIII**, **XII** or **XVI** in target validation. Target validation is the disruption of an identified DNA sequence to ascertain the function of the sequence, and a compound of formula **II**, **VIII**, **XII** or **XVI** can be used to selectively bind an identified sequence, and thus disrupt its function, i.e. functional genomics

Preferred Synthetic Strategies

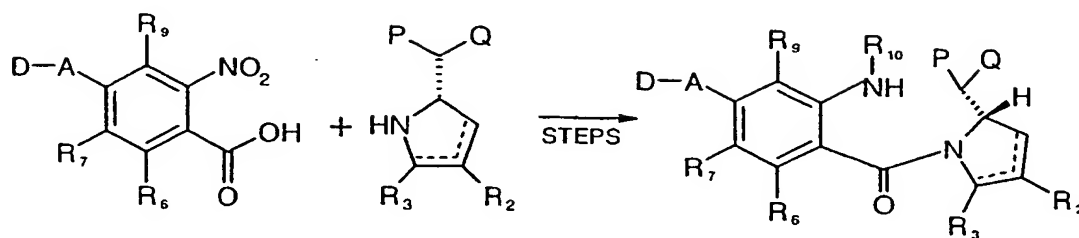
A key step in a preferred route to compounds of formula **I** is a cyclisation process to produce the B-ring, involving generation of an aldehyde (or functional equivalent thereof) at what will be the 11-position, and attack thereon by the pro-10-nitrogen:



In this structure, D represents XY, or a masked form thereof. The "masked aldehyde" -CPQ may be an acetal or thioacetal, in which case the cyclisation involves unmasking. Alternatively, the masked aldehyde may be an aldehyde precursor, such as an alcohol -CHOH, in which case the reaction involves oxidation, e.g. by means of TPAP or DMSO (Swern oxidation).

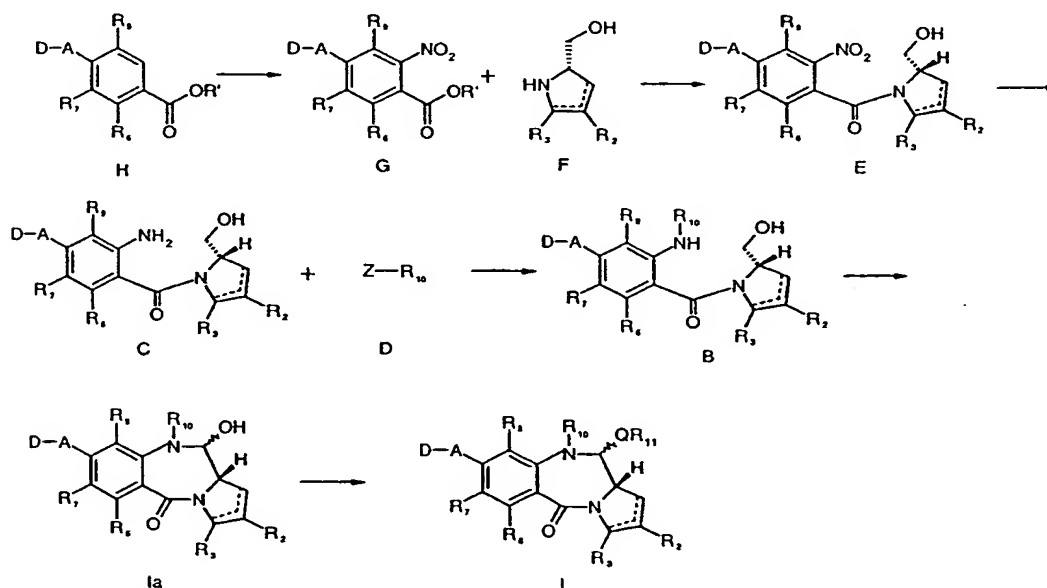
The masked aldehyde compound can be produced by condensing a

corresponding 2-substituted pyrrolidine with a 2-nitrobenzoic acid:



The nitro group can then be reduced to $-NH_2$, and protected by reaction with a suitable agent, e.g. a chloroformate, which provides the removable nitrogen protecting group in the compound of formula I.

A process involving the oxidation-cyclization procedure is illustrated in scheme 1 (an alternative type of cyclisation will be described later with reference to scheme 2).



Scheme 1

If R_{11} is other than hydrogen, the compound of formula I, may be prepared by direct etherification of the alcohol Ia.

Compounds with $Q = S$ can be prepared by treatment of the corresponding alcohol Ia with $R_{11}SH$, and a catalyst (usually an

acidic solution HCl, and sometimes a Lewis Acid such as Al_2O_3). If $Q = NH$, then these compounds can be prepared by reacting Ia with an amine $R_{11}NH$ and a catalyst (usually an aqueous acid or a Lewis Acid).

Exposure of the alcohol B (in which the 10-nitrogen is generally protected as a carbamate) to tetrapropylammonium perruthenate (TPAP)/N-methylmorpholine N-oxide (NMO) over A4 sieves results in oxidation accompanied by spontaneous B-ring closure to afford the desired product. The TPAP/NMO oxidation procedure is found to be particularly convenient for small scale reactions while the use of DMSO-based oxidation methods, particularly Swern oxidation, proves superior for larger scale work (e.g. > 1 g).

The uncyclized alcohol B may be prepared by the addition of a nitrogen protecting reagent of formula D, which is preferably a chloroformate or acid chloride, to a solution of the amino alcohol C, generally in solution, generally in the presence of a base such as pyridine (preferably 2 equivalents) at a moderate temperature (e.g. at $0^{\circ}C$). Under these conditions little or no O-acylation is usually observed.

The key amino alcohol C may be prepared by reduction of the corresponding nitro compound E, by choosing a method which will leave the rest of the molecule intact. Treatment of E with tin (II) chloride in a suitable solvent, e.g. refluxing methanol, generally affords, after the removal of the tin salts, the desired product in high yield.

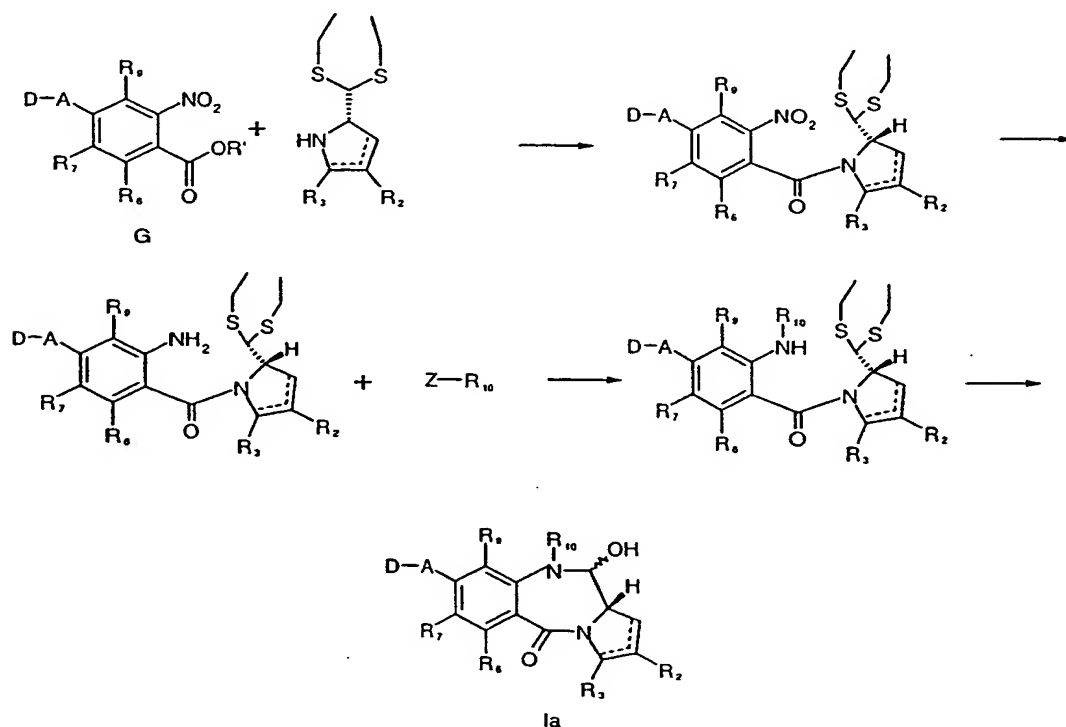
Exposure of E to hydrazine/Raney nickel avoids the production of tin salts and may result in a higher yield of C, although this method is less compatible with the range of possible C and A-ring substituents. For instance, if there is C-ring unsaturation (either in the ring itself, or in R_2 or R_3), this technique may be unsuitable.

The nitro compound of formula E may be prepared by coupling

the appropriate *o*-nitrobenzoyl chloride to a compound of formula **F**, e.g. in the presence of K_2CO_3 at $-25^\circ C$ under a N_2 atmosphere. Compounds of formula **F** can be readily prepared, for example by olefination of the ketone derived from L-trans-4-hydroxy proline. The ketone intermediate can also be exploited by conversion to the enol triflate for use in palladium mediated coupling reactions.

The *o*-nitrobenzoyl chloride is synthesised from the *o*-nitrobenzoic acid (or, after hydrolysis, the alkyl ester) of formula **G**, which itself is prepared from the vanillic acid (or alkyl ester) derivative **H**. Many of these are commercially available and some are disclosed in Althuis, T. H. and Hess, H. J., *J. Medicinal Chem*, 20(1), 146-266 (1977).

Alternative Cyclisation (Scheme 2)



Scheme 2

In scheme 1, the final or penultimate step was an oxidative cyclisation. An alternative approach, using thioacetal coupling, is shown in scheme 2. Mercury-mediated unmasking

causes cyclisation to the desired compound (Ia).

The thioacetal intermediates may be prepared as shown in scheme 2: the thioacetal protected C-ring [prepared via a literature method: Langley, D.R. & Thurston, D.E., *J. Organic Chemistry*, 52, 91-97 (1987)] is coupled to the o-nitrobenzoic acid (or, after hydrolysis, the alkyl ester) G using a literature procedure. The resulting nitro compound cannot be reduced by hydrogenation, because of the thioacetal group, so the tin (II) chloride method is used to afford the amine. This is then N-protected, e.g., by reaction with a chloroformate or acid chloride, such as p-nitrobenzylchloroformate.

Acetal-containing C-rings can be used as an alternative in this type of route with deprotection involving other methods including the use of Lewis Acid conditions.

In the above synthesis schemes, the derivatisation of the A-ring is shown as being complete before the compounds are attached to the solid support. This is preferred if the substituents are groups such as alkoxy or nitro. On the other hand, substituent groups such as alkyl or alkenyl could be added to the A-ring after the coupling of the compound to the solid support. This may be achieved by R₆, R₇, or R₈ being easily replaceable groups, such as halogen atoms.

An alternative synthesis approach to those detailed above is to protect the pro N10 position on the component which will form the A-ring, before joining the component which will form the C-ring.

Embodiments of the present invention will now be described by way of example with reference to the accompanying drawings in which:

Fig. 1 is a reaction scheme for the synthesis of compounds of formula I;

Fig. 2 is a reaction scheme for the synthesis of alternative compounds of fomula I;

Figs. 3a and 3b are a reaction scheme for the synthesis of compounds of formulae V, IV, III, and II;

5 Figs. 4a and 4b are a reaction scheme for the synthesis of further compounds of formulae V, IV, III, and II;

Fig. 5 is a reaction scheme for the synthesis of compounds of formula III and IV;

10 Figs. 6a, 6b and 6c are a reaction scheme for the synthesis of compounds of formula VI and VII;

Fig. 7 is a reaction scheme for the synthesis of compounds of formula III and IV;

Figs. 8 and 9 are reaction schemes for the synthesis of compounds of formula III and IV;

15 Figs. 10a,b and c are a reaction scheme for the synthesis of compounds of formula X and XI;

Fig. 11 is reaction schemes for the synthesis of compounds of formula XIII and XIV; and

20 Figs. 12 to 15 are reaction schemes for the synthesis of compounds of formula III and IV.

General Methods

Melting points (mp) were determined on an Electrothermal 9100 digital melting point apparatus and are uncorrected. Infrared
25 (IR) spectra were recorded using a Perkin-Elmer Spectrum 1000 spectrophotometer. ¹H- and ¹³C- NMR spectra were recorded on a Jeol GSX 270 MHz FT-NMR spectrometer operating at 20°C +/-1°C. Chemical shifts are reported in parts per million (δ) downfield from tetramethylsilane (TMS). Spin multiplicities
30 are described as: s (singlet), bs (broad singlet), d (doublet), dd (doublet of doublets), t (triplet), q (quartet), p (pentuplet) or m (multiplet). Mass spectra (MS) were recorded using a Jeol JMS-DX 303 GC Mass Spectrometer (EI mode: 70eV, source 117-147°C). Accurate molecular masses
35 (HRMS) were determined by peak matching using perfluorokerosene (PFK) as an internal mass marker, and FAB mass spectra were obtained from a glycerol/thioglycerol/trifluoroacetic acid (1:1:0.1) matrix

with a source temperature of 180°C. Optical rotations at the Na-D line were obtained at ambient temperature using an ADP 220 Automatic Polarimeter (Bellingham & Stanley). Flash chromatography was performed using Aldrich flash chromatography "Silica Gel-60" (E. Merck, 230-400 mesh). Thin-layer chromatography (TLC) was performed using GF₂₅₄ silica gel (with fluorescent indicator) on glass plates. All solvents and reagents, unless otherwise stated, were supplied by the Aldrich Chemical Company Ltd. and were used as supplied without further purification. Anhydrous solvents were prepared by distillation under a dry nitrogen atmosphere in the presence of an appropriate drying agent, and were stored over 4Å molecular sieves or sodium wire. Petroleum ether refers to the fraction boiling at 40-60°C.

Example 1: Synthesis of PBDs of formula I (Figure 1)

Overall Synthesis

The compounds with an acid-terminating side chain, 7a-c (R_{10} = Nvoc, Fmoc, Teoc), were prepared by palladium-mediated de-esterification of the appropriate allyl esters. The esters were in turn prepared by Swern oxidation (oxidation of the primary alcohol to an aldehyde which provokes spontaneous B-ring closure) of the Nvoc, Fmoc and Teoc protected amino alcohols. The carbamate-protected amino alcohols were prepared by treating the common amino alcohol intermediate 4 with the appropriate chloroformate in the presence of pyridine. The amino alcohol was obtained by reduction of the nitro compound 3, which in turn was assembled by coupling pyrrolidine methanol to the o-nitrobenzoic acid 2. Compound 2 was prepared by selective esterification of the diacid 1 at the aliphatic acid. Finally, the diacid was obtained by simultaneous nitration and oxidation of the known hydroxypropyloxy vanillic acid derivative 0.

The Troc-protected compound 7d was prepared by an alternative synthetic strategy involving the use of acetals. The ring closed allyl ester 6d was prepared by unmasking the acetal protected aldehyde in the presence of the Troc protected

amine. The Troc protected amine was obtained through exposure of the free amine 9 to Troc-Cl in the presence of pyridine. Reduction with tin chloride furnished the amine 9 from the nitro acetal 8, which in turn was obtained by coupling 2 to the appropriate acetal-protected prolinal.

Allyl Amino Alcohol Intermediate (4)

3-(4-carboxy-2-methoxy-5-nitrophenoxy)propanoic acid (1)

The alcohol 0 (50 g, 0.22 mol) was added portionwise over 1 hour to nitric acid (70%, 400 ml) cooled to 0°C. Once addition was complete, the solution was stirred at 0°C for 1 hour, then allowed to warm to RT. The semisolid formed was collected by filtration and washed with a minimum of ice/water. The resulting pale yellow solid was redissolved in EtOAc, the solution dried (MgSO₄) and then concentrated to afford the diacid 1 (31 g, 49%). ¹H NMR (270 MHz): δ 2.83-2.79 (t, J = 6, 12.5 Hz, 2H), 3.94 (s, 3H), 4.37-4.33 (t, J = 6, 12.5 MHz, 2H), 7.18 (s, 1H), 7.46 (s, 1H), 10.38 (br.s, 2H).

2-Propene 3-(4-carboxy-2-methoxy-5-nitrophenoxy)propanoate (2)

A mixture of 3-(4-carboxy-2-methoxy-5-nitrophenoxy)propanoic acid 1 (20 g, 74.3 mmol) and p-toluene sulphonic acid monohydrate (2.3 g, 7.4 mmol) in allyl alcohol (240 mL, 3.5 mol) was refluxed for 7 hours then allowed to cool. The allyl alcohol was then removed in vacuo, and the residue triturated with dilute HCl acid (3 x 75 ml) and collected by filtration. This solid was taken up in EtOAc, and the resulting solution washed with water (3 x 50 ml) and brine (3 x 50 ml) and dried over sodium sulphate. Evaporation in vacuo afforded 2 as a white solid (19.27 g, 84%): mp 128-130°C; ¹H-NMR (270 MHz, CDCl₃) δ 2.92 (t, 2H, J = 6.35 Hz); 3.94 (s, 3H); 4.38 (t, 2H, J = 6.41 Hz); 4.65 (d, 2H, J = 5.61 Hz); 5.27 (dd, 1H, J₁ = 1.28 Hz, J₂ = 19.42 Hz); 5.33 (dd, 1H, J₁ = 1.28 Hz, J₂ = 17.04 Hz); 5.92 (m, 1H); 7.15 (s, 1H); 7.45 (s, 1H); ¹³C NMR (67.8 MHz, CDCl₃): δ 34.1, 56.5, 65.0, 65.4, 108.5, 111.3, 118.3, 122.9, 131.8, 141.1, 149.1, 152.6, 167.1, 170.0; IR (Nujol); ν 1730, 1630, 1550, 1430, 1390, 1290, 1230, 1190, 1170, 1070, 1030, 1010 cm⁻¹; MS (EI) m/z (relative intensity): 325 (M⁺,

19), 251 (3), 213 (2), 196 (3), 211 (3), 113 (19), 91 (4), 71 (9), 55 (6); HRMS: calcd. for $C_{14}H_{15}NO_8$ 325.0798, found 232.0773.

2-Propene 3-(4-[2'-hydroxymethylpyrrolidinecarboxy]-2-methoxy-5-nitrophenoxy)propanoate (3)

Oxalyl chloride (2.7 ml, 31.0 mmol) was added dropwise to a suspension of the nitro acid 2 (9 g, 28.0 mmol) and DMF (0.05 ml) in CH_2Cl_2 (150 ml), followed by stirring at room temperature for 16 hours. The resulting solution was added dropwise to a stirred solution of pyrrolidine methanol (3 ml, 31.0 mmol) and triethylamine (8.5 ml, 61.0 mmol) in CH_2Cl_2 (80 ml) at $-20^\circ C$ (liquid N_2 /acetone) followed by stirring at room temperature for 16 hours under N_2 . After quenching with aqueous HCl (1.0 N, 50 ml), the separated organic phase was washed with H_2O (3 x 25 ml) and brine (3 x 10 ml), dried over magnesium sulphate and evaporated *in vacuo* to afford a crude orange oil. Purification by flash column chromatography (5% MeOH/EtOAc) afforded 3 as a pale yellow oil (7.9 g, 70%): 1H NMR (270 MHz, $CDCl_3$): δ 2.22-1.71 (m, 6H); 2.94 (t, J = 6.4 Hz, 2H); 3.15 (d x d, J = 6.5 Hz, 2H); 3.92-3.76 (m, 1H); 3.96 (s, 3H); 4.4 (t, J = 6.2 Hz, 2H); 4.67-4.64 (m, 2H); 5.39-5.23 (m, 2H); 6.0-5.86 (m, 1H); 6.81 (s, 1H); 7.75 (s, 1H); ^{13}C NMR (67.8 MHz, $CDCl_3$): δ 24.4, 28.5, 34.1, 49.5, 56.7, 61.6, 64.9, 65.6, 108.9, 109.3, 118.6, 128.2, 131.8, 148.2, 154.9, 170.1; IR (film): ν 3394, 2947, 2882, 1735, 1689, 1618, 1577, 1521, 1454, 1431, 1386, 1334, 1276, 1221, 1178, 1059, 1002 cm^{-1} ; MS (EI) M/Z (relative intensity): 408 (M^+ , 1), 390(4), 377(20), 308(86), 296(3), 278(9), 265(35), 252(3), 118(74), 111(3), 108(8), 98(4), 83(14). HRMS: calcd. for $C_{19}H_{23}O_6N_2$ 377.416, found 377.1711.

2-Propene 3-(5-amino-4-[2'-hydroxymethylpyrrolidinecarboxy]-2-methoxyphenoxy)propanoate (4)

Solid $SnCl_4 \cdot 2H_2O$ (21.3 g, 0.095 mol) was added to a stirred solution of the nitro alcohol 3 (7.7 g, 0.02 mol) in MeOH (100 ml), and the mixture heated at reflux for 45 min. The solvent was then evaporated *in vacuo*, and the residual oil partitioned between EtOAc (50 ml) and aqueous saturated $NaHCO_3$ (50 ml).

followed by vigorous stirring for 16 hours to aid separation. The combined layers were filtered through Celite and washed with EtOAc (25 ml). The layers were separated and the resulting organic phase was washed with H₂O (3 x 25 ml) and brine (3 x 10 ml) and then dried over magnesium sulphate. Evaporation *in vacuo* afforded the amine **4** as a dark orange oil (5.6 g, 78%): ¹H NMR (270 MHz, CDCl₃) δ 2.17-1.65 (m, 6H); 2.9 (t, *J* = 6.6 Hz, 2H); 3.72-3.46 (m, 3H); 3.75 (s, 3H); 4.2 (t, *J* = 6.8 Hz, 2H); 4.4 (br. d x d, *J* = 9.7 Hz, 2H); 4.65-4.62 (m, 2H); 5.37-5.22 (m, 2H); 6.0-5.85 (m, 1H); 6.3 (s, 1H); 6.76 (s, 1H). ¹³C NMR (67.8 MHz, CDCl₃): δ 24.9, 28.7, 34.3, 57.3, 61.2, 64.2, 65.5, 67.4, 102.6, 113.5, 118.5, 131.9, 141.1, 150.9, 170.5. IR (film): ν 3354, 2940, 2880, 1734, 1621, 1589, 1514, 1453, 1429, 1407, 1265, 1230, 1173, 1110, 1023. MS(EI) *M/z* (relative intensity): 378 (*M*⁺, 60), 278(100), 266(5), 252(9), 238(3), 220(4), 206(6), 194(4), 178(3), 166(40), 150(5), 137(20), 123(4), 113(4), 107(4), 100(8), 94(12), 84(9). HRMS: calcd. for C₁₁H₁₆O₆N₂ 378.424, found 378.1760.

Example 1(a): Nvoc-PBD acid (7a)

Nvoc Chloroformate

4,5-Dimethoxynitrobenzyl alcohol (2 g, 9.4 mmol) and triphosgene (0.93 g, 3.13 mmol) were dissolved in CH₂Cl₂ (50 ml), and the resulting red suspension stirred vigorously and cooled to 0°C. Pyridine (260 μl, 3.13 mmol) was added dropwise, and the resulting green solution stirred at room temperature for 16 hours to afford a solution of the chloroformate that was used directly in the next step.

Allyl Nvoc Alcohol (5a)

A solution of freshly prepared (see above) Nvoc chloroformate (9.4 mmol) and pyridine (1.3 ml, 9.5 mmol) was added dropwise to a stirred solution of the amino alcohol **4** (3 g, 7.9 mmol) in CH₂Cl₂ (60 ml) at 0°C. The mixture was allowed to return to room temperature and stirring continued for 3 hours. Evaporation *in vacuo* afforded an oil which was redissolved in CH₂Cl₂ (50 ml). The resulting solution was washed with HCl (1.0 N, 3 x 25 ml), H₂O (3 x 25 ml) and brine (3 x 10 ml), dried

over magnesium sulphate and evaporated *in vacuo* to give a dark yellow foam. This was purified by flash column chromatography (EtOAc) to afford the carbamate 5a as a pale yellow foam (3.7 g, 76%): ¹H NMR (270 MHz, CDCl₃): δ 1.6-2.2 (m, 6H); 2.9 (t, *J* = 6.2, 2H); 3.4-3.9 (m, 3H); 3.81 (s, 3H); 3.97 and 4.01 (2 x s, 6H); 4.3 (t, *J* = 6.4 Hz, 2H); 4.63 (d, *J* = 5.9 Hz, 2H); 5.22-5.36 (m, 2H); 5.5-5.67 (m, 2H); 5.85-6.0 (m, 1H); 6.84 (s, 1H); 7.09 (s, 1H); 7.74 (s, 1H); 8.93 (br. s, 1H). ¹³C NMR (67.8MHz, CDCl₃): δ 25.1, 28.3, 34.3, 51.5, 56.4, 56.6, 56.8, 61.0, 63.8, 64.4, 65.4, 66.4, 106.3, 108.2, 110.0, 111.9, 118.5, 127.8, 131.5, 131.9, 139.6, 144.4, 148.1, 150.3, 153.2, 153.7, 170.3, 170.7. IR (reflectance): ν 3329, 3110, 2937, 1728, 1581, 1523, 1453, 1323, 1270, 1175, 1129, 1070, 1031, 1011. MS(FAB) *m/z* (relative intensity): 618 (*M*⁺+1(3)), 473(2), 439(1), 405(1), 378(3), 304(7), 278(16), 196(100), 166(28), 151(15), 102(27), 70(9).

Allyl Nvoc PBD (6a)

A solution of DMSO (1.45 ml, 0.02 mol) in CH₂Cl₂ (40 ml) was added over 45 minutes to a stirring solution of oxalyl chloride (5.1 ml, 0.01 mol) in CH₂Cl₂ (20 ml) cooled to -40°C (liquid N₂/chlorobenzene). Stirring was continued for a further 15 minutes at -40°C, and then a solution of the NVOC alcohol 5a (3.5 g, 5.7 mmol) in CH₂Cl₂ (45 ml) was added dropwise over 1 hour. Stirring was continued at -40°C for a further 45 min, and then a solution of Et₃N (3.4 ml, 0.024 mol) in CH₂Cl₂ (20 ml) was added dropwise over 30 minutes and stirring continued for 1 hour. The mixture was then allowed to warm to room temperature before diluting with CH₂Cl₂ (20 ml). The organic phase was washed with HCl (1.0 N) (3 x 50 ml), H₂O (3 x 50 ml) and brine (3 x 25 ml), dried over magnesium sulphate and then evaporated *in vacuo* to give a yellow foam. This was purified by flash column chromatography (1% MeOH/CHCl₃) to afford 6a as a pale yellow foam (3.2 g, 91%): ¹H NMR (270 MHz, CDCl₃): δ 1.9-2.2 (m, 6H); 2.86 (t, *J* = 6.9 Hz, 2H); 3.45-3.6 (m, 3H); 3.81 (s, 3H); 3.88 and 3.91 (2 x s, 6H); 4.2-4.4 (m, 3H); 4.61 (m, 2H); 5.19-5.35 (m, 2H); 5.49 (s, 2H); 5.7 (br d, *J* = 9.9 Hz, 1H); 5.82-5.97 (m, 1H); 6.51

(s, 1H); 6.86 (s, 1H); 7.25 (s, 1H); 7.66 (s, 1H); ¹³C NMR (67.8 MHz, CDCl₃): δ 23.1, 28.7, 30.6, 34.1, 46.5, 56.2, 60.1, 64.6, 65.3, 65.5, 86.1, 107.9, 109.2, 110.8, 114.3, 118.4, 126.7, 127.0, 128.1, 131.8, 138.9, 147.9, 148.9, 149.9, 153.8, 155.4, 166.8, 170.3. IR (reflectance): ν 3329, 3084, 2940, 1713, 1633, 1519, 1454, 1276, 1105, 1067. MS (EI) M/Z (relative intensity) 615 (M⁺, 12), 503(100), 358(4), 261(3), 246(37), 231(4), 196(32), 180(24), 166(4), 150(4), 136(6), 70(31).

Acid Nvoc PBD (7a)

Tetrakis(triphenylphosphine) palladium (0.583 g, 0.504 mmol) and morpholine (4.4 ml, 50.4 mmol) were added to a solution of the allyl carbinolamine 6a (3.1 g, 5.04 mmol) in THF (30 ml) and the mixture stirred for 16 hours. After evaporation in vacuo, the resulting oil was redissolved in CH₂Cl₂, and the solution washed with HCl (1.0 N) (3 x 25 ml), H₂O (3 x 25 ml) and brine (3 x 10 ml), dried over magnesium sulphate and then evaporated in vacuo to give an orange foam. This was purified by flash column chromatography (1% MeOH/CHCl₃) to afford 7a as a pale yellow foam (2.25 g, 78%): ¹H NMR (270 MHz, CDCl₃): δ 1.9-2.3 (m, 4H); 2.85 (t, J = 6.7 Hz, 2H); 3.4-3.6 (m, 3H); 3.68 (s, 3H); 3.87 and 3.9 (2 x s, 6H); 4.2-4.4 (m, 3H); 5.4-5.48 (m, 2H); 5.7 (br d, J = 9.9 Hz, 1H); 6.5 (s, 1H); 6.89 (s, 1H); 7.26 (s, 1H); 7.64 (s, 1H); ¹³C NMR (67.8 MHz, CDCl₃): δ 23.0, 28.5, 33.9, 46.6, 56.2, 60.4, 64.6, 65.4, 86.1, 107.9, 109.3, 110.8, 114.7, 126.5, 126.9, 128.2, 138.9, 147.9, 148.9, 149.9, 153.8, 155.5, 167.1, 174.5. IR (reflectance): ν 2939, 2252, 1712, 1599, 1522, 1459, 1277, 1221, 1137, 1105, 1066, MS (FAB) M/Z (relative intensity): 576 (M⁺+1, 15), 514(3), 381(3), 363(3), 336(3), 319(9), 303(2), 293(3), 289(2), 279(4), 266(6), 264(14), 253(3), 245(4), 238(10), 215(4), 206(4), 196(100), 192(16), 180(23), 166(32), 151(18), 136(10), 123(7), 117(14), 93(28), 73(15), 70(20).

Example 1(b): Fmoc-PBD acid (7b)

Allyl Fmoc Alcohol (5b)

9-Fluorenylmethyl chloroformate (5.65 g, 0.022 mol) was added

portionwise to a stirred solution of the amino alcohol **4** (7.5 g, 0.02 mol) and Na_2CO_3 (5.26 g, 0.05 mol) in a mixture of THF (150 ml) and water H_2O (150 ml) at 0°C . The reaction mixture was allowed to return to room temperature, stirred for a further 2 h, and then extracted with EtOAc (3 x 50 ml). The combined organic phase was washed with H_2O (3 x 50 ml) and brine (3 x 25 ml), dried over magnesium sulphate and evaporated *in vacuo* to give a dark red oil. This was purified by flash column chromatography (petroleum ether 40-60/EtOAc, 1:1) to afford **5b** as a pale yellow oil (8.11 g, 68%): ^1H NMR (270 MHz, CDCl_3): δ 1.72-2.18 (m, 6H); 2.9 (t, $J = 6.4$, 2H); 3.43-3.91 (m, 3H); 3.81 (s, 3H); 4.25-4.54 (m, 5H); 4.61-4.65 (m, 2H); 5.21-5.37 (m, 2H); 5.85-6.0 (m, 1H); 6.85 (s, 1H); 7.31-7.79 (m, 9H); 8.77 (br s, 1H); ^{13}C NMR (67.8 MHz, CDCl_3): δ 25.1, 28.4, 35.3, 47.0, 56.7, 60.9, 64.3, 65.4, 66.3, 67.1, 106.4, 111.7, 118.3, 120.0, 125.2, 127.1, 127.2, 127.8, 131.5, 131.9, 141.3, 143.7, 144.4, 150.2, 153.8, 170.3; MS (FAB): 601 ($\text{M}^+ + 1$); HRMS: calcd for $\text{C}_{34}\text{H}_{36}\text{N}_2\text{O}_8$ 600.667, found 600.2175. IR (film): ν 3315, 2952, 1727, 1597, 1522, 1452, 1392, 1322, 1174, 1117, 1017.

Fmoc Allyl Carbinolamine Cyclised (**6b**)

A solution of DMSO (3.4 ml, 0.048 mol) in CH_2Cl_2 (100 ml) was added dropwise over 45 minutes to a stirred solution of oxalyl chloride (12 ml, 0.024 mol) in CH_2Cl_2 (50 ml) at -40°C (liquid N_2 /chlorobenzene). The mixture was stirred at -40°C for a further 15 minutes and then a solution of the Fmoc alcohol **5b** (8g, 0.013 mol) in CH_2Cl_2 (135 ml) was added over 1 hour. After stirring for a further 45 minutes at -40°C , a solution of DIPEA (10 ml, 0.057 mol) in CH_2Cl_2 (55 ml) was added over 30 minutes and stirring continued for 1 hour. The solution was then allowed to warm to room temperature, diluted with CH_2Cl_2 (100 ml), and the organic phase washed with HCl (1.0 N) (3 x 50 ml), H_2O (3 x 50 ml) and brine (3 x 25 ml), dried over magnesium sulphate and evaporated *in vacuo* to afford **6b** as a pale cream foam (6.4 g, 80%): ^1H NMR (270 MHz, CDCl_3): δ 1.99-2.1 (m, 4H); 2.83-2.87 (m, 2H); 3.51-3.6 (m, 2H); 3.6-3.8 (m, 1H); 3.95 (s, 3H); 4.0-4.58 (m, 7H); 5.17-5.31 (m, 2H); 5.68

(d, $J = 9.7$ Hz, 1H); 5.84-5.87 (m, 1H); 6.75 (s, 1H); 7.02-7.75 (m, 9H); ^{13}C NMR (67.8 MHz, CDCl_3) δ 23.0, 28.7, 34.2, 46.5, 53.5, 56.2, 59.5, 60.1, 64.4, 65.4, 68.4, 86.0, 111.2, 114.7, 118.4, 119.9, 124.9, 125.4, 126.8, 127.1, 127.8, 128.3, 131.8, 131.9, 141.1, 141.2, 143.1, 143.5, 148.9, 149.9, 156.1, 166.9, 170.3; MS (FAB) 599 ($\text{M}^+ + 1$). IR (reflectance): ν 3318, 2950, 1713, 1603, 1517, 1386, 1290, 1177, 1037.

Fmoc Acid Carbinolamine (7b)

Phenylsilane (2.5 ml, 0.02 mol) and tetrakis (triphenylphosphine) palladium (0.232 g, 0.2 mmol) were added to a solution of the Fmoc carbinolamine **6b** (6 g, 0.01 mol) in CH_2Cl_2 (80 ml) followed by stirring at room temperature for 16 hours. The reaction was quenched with H_2O (50 ml) and extracted with CH_2Cl_2 (3 x 30 ml). The combined organic phase was washed with water (3 x 30 ml), brine (3 x 25 ml), dried over magnesium sulphate and evaporated *in vacuo* to give a dark brown foam. This was purified by flash column chromatography ($\text{MeOH}/\text{CHCl}_3$, 1:99) to afford **7b** as a pale beige foam (4.3 g, 77%): ^1H NMR (270 MHz, CDCl_3): δ 1.9-2.2 (m, 4H); 2.65-2.85 (m, 2H); 3.4-3.6 (m, 2H); 3.6-3.8 (m, 1H); 3.91 (s, 3H); 4.0-4.25 (m, 4H); 4.45-4.5 (m, 1H); 5.68 (d, $J = 9.5$ Hz, 1H); 6.75 (s, 1H); 6.9-7.7 (m, 9H); ^{13}C NMR (67.8 MHz, CDCl_3) δ 23.0, 28.6, 33.9, 46.5, 56.2, 60.3, 64.6, 68.5, 86.0, 111.2, 115.2, 119.8, 124.9, 126.8, 127.1, 127.7, 128.2, 140.9, 141.1, 142.9, 143.4, 149.1, 149.9, 156.3, 167.0, 174.6. IR (reflectance); ν 3316, 2955, 2609, 2249, 1713, 1601, 1514, 1453, 1279, 1036. MS (FAB) M/Z (relative intensity) 560 ($\text{M}^+ + 1$).

Example 1(c): Teoc-PBD Acid (7c)

Allyl Teoc Alcohol (5c)

Pyridine (0.165 ml, 2.04 mmol) was added dropwise to a solution of triphosgene (0.605g, 2.04 mmol) and 2-trimethylsilyl ethanol (1.082 g, 9.15 mol) in anhydrous CH_2Cl_2 (100 ml), and the mixture allowed to stir at room temperature for 16 hours. This solution was added dropwise to a stirred solution of the amino alcohol **4** (2.30 g, 6.10 mmol) and pyridine (0.987 mL, 0.0122 mol) in anhydrous CH_2Cl_2 (50 mL) at

0°C (ice bath) under a nitrogen atmosphere. After reaction was complete as indicated by TLC (petroleum ether/ethyl acetate, 1:1), the mixture was washed with copper (II) sulphate (2 x 100 mL) and brine (100 mL), dried (MgSO₄) and evaporated in vacuo to give a brown oil. This was purified by flash column chromatography (chloroform/methanol, 99:1) to afford **5c** as a brown solid (2.5 g, 78%): ¹H NMR (270 MHz, CDCl₃): δ -0.06 (s, 9H), 1.01 (m, 2H), 1.82-2.30 (m, 4H), 2.86 (m, 2H), 3.40-3.75 (m, 7H), 4.15-4.31 (m, 4H), 4.6 (m, 2H), 5.15-5.31 (m, 2H), 5.80-5.94 (m, 1H), 6.76 (s, 1H), 7.76 (s, 1H), 8.52 (s, 1H); ¹³C NMR (67.8 MHz, CDCl₃) δ -1.47, 17.7, 25.1, 28.4, 34.3, 51.6, 56.8, 61.2, 63.5, 64.3, 65.4, 66.8, 106.2, 112.1, 113.8, 118.3, 132.0, 132.2, 144.0, 154.1, 170.3; MS (EI): 522 (M⁺, 12.8), 435 (7), 350 (13), 319 (77), 262 (27), 206 (13), 149 (88), 83 (32), 70 (100); HRMS: Calcd 522.2397, found 522.2351.

Allyl Teoc Carbinolamine PBD (**6c**)

A solution of DMSO (1.02 mL, 0.014 mol) in dry CH₂Cl₂ (30 mL) was added to a solution of oxalyl chloride (3.59 mL, 7.185 mmol) in CH₂Cl₂ (25 mL) at -43°C (chlorobenzene/liq. N₂) under a nitrogen atmosphere. After stirring at -43°C for 45 min, a solution of the TEOC alcohol **5c** (2.50 g, 4.79 mmol) in dry CH₂Cl₂ (30 mL) was added dropwise to the reaction mixture and stirring continued at -43°C for a further 45 min. A solution of triethylamine (3.34 mL, 0.024 mol) in dry DCM (25 mL) was then added dropwise, and the vessel allowed to warm to 0°C. The reaction mixture was diluted with CH₂Cl₂ (150 mL), washed with 1N HCl (100 mL), water (100 mL) and brine (100 mL), dried (MgSO₄) and then evaporated in vacuo to give crude **6c**. This was purified by flash column chromatography (silica gel, chloroform) to afford **6c** as a yellow oil (1.72g, 69%): ¹H NMR (270 MHz, CDCl₃): δ -0.08 (s, 9H), 0.92 (m, 2H), 2.04-2.33 (m, 4H), 3.14 (m, 2H), 3.50-3.75 (m, 4H), 3.93 (s, 3H), 4.00-4.40 (m, 4H), 4.67 (m, 2H), 5.26-5.40 (m, 2H), 5.65 (d, 1H, J = 9.52 Hz), 5.89-5.99 (m, 1H), 6.72 (bs, 1H), 7.23 (s, 1H); ¹³C NMR (67.8 MHz, CDCl₃) δ -1.47, 17.6, 23.0, 28.7, 34.2, 46.4, 56.2, 59.9, 64.3, 65.4, 65.5, 85.9, 111.0, 114.7, 118.3, 126.4, 131.8, 132.0, 148.7, 149.7, 154.2, 170.0, 170.4; MS

(FAB): 629 (0.8), 593 (0.91), 536 (1.5), 493 (4.6), 465 (1.0), 449 (1.7), 431 (6.8), 394 (8.1), 368 (1.3), 338 (1.5), 304 (5.8), 264 (3.6), 238 (2.2), 204 (1.6), 192 (9.1), 166 (2.5), 149 (6.8), 98 (4.3), 73 (100).

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Acid Teoc PBD Carbinolamine (7c)

Tetrakis(triphenylphosphine)palladium(0) (190 mg, 0.165 mmol) was added to a solution of the Teoc-protected carbinolamine **6c** (1.72 g, 3.30 mmol) in ethanol (50 mL), and the mixture heated at reflux for 60 minutes after which time TLC (AcOH/MeOH/chloroform, 1:10:100) indicated that reaction was complete. The reaction mixture was allowed to cool and was then filtered through Celite. Evaporation of the solvent in vacuo afforded **7c** as a yellow solid (1.08g, 68%): ¹H NMR (270 MHz, CDCl₃): δ -0.06 (s, 9H), 0.86 (m, 2H), 1.98-2.20 (m, 4H), 2.8-3.0 (m, 2H), 3.40-3.70 (m, 3H), 3.75 (s, 3H), 4.00-4.40 (m, 2H), 5.65 (d, J = 8.63 Hz, 1H), 6.78 (bs, 1H), 7.21 (s, 1H): ¹³C NMR (67.8 MHz, CDCl₃) δ, -1.5, 18.3, 23.1, 28.7, 34.5, 46.4, 56.1, 58.4, 64.8, 64.9, 85.9, 110.8, 115.0, 126.3, 128.7, 148.6, 149.6, 167.2.

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20**Example 1(d): Synthesis of Troc-PBD acid 7d****Prop-2-enyl 4-(N-2S-Diethylthiomethylpyrrolidinecarboxy)-2-methoxy-5-nitrophenyl propanoate (8)**

2-Propene 3-(4-carboxy-2-methoxy-5-nitrophenyloxy)propanoate **2**: 5 g, 15.34 mmol), oxalyl chloride (2 mL, 23 mmol) and 5 drops of DMF were stirred in dry THF (100 mL) for 18 hours. The solvent was then removed in vacuo and the residue dissolved in dry THF (50 mL). This was added dropwise to a vigorously stirred mixture of (2S)-pyrrolidine-2-carboxaldehyde diethyl thioacetal (3.15 g, 15.34 mmol) and triethylamine (1.86 g, 18.41 mmol). The stirring was continued for 18 hours. The solvent was then removed in vacuo and the product purified by flash column chromatography (ethyl acetate) to give **8** (7.48g, 95%) as a yellow oil. ¹H NMR (270 MHz, CDCl₃): δ 7.74 (s, 1H, OCCHC), 6.83 (s, 1H, MeOCCHC), 5.98-5.86 (m, 1H, CH₂CHCH₂), 5.33 (d, 1H, J = 26.56 Hz, OCH₂CHCH₂), 5.28 (d, 1H, J = 20.24 Hz, OCH₂CHCH₂), 4.88 (d, 1H,

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J = 3.85 Hz, NCHCH), 4.74-4.65 (m, 2H, OCH₂CHCH₂) 4.42 (t, 2H, J = 7.69 Hz, CH₂CH₂OC), 3.94 (s, 3H, OCH₃), 3.29-3.21 (m, 2H, NCH₂), 2.96 (p, 2H, J = 3.12 Hz, CH₂CH₂O), 2.87-2.67 (m, 4H, SCH₂CH₃), 2.32-1.78 (m, 4H, NCH₂CH₂CH₂) 1.38-1.31 (m, 6H, SCH₂CH₃).

¹³C-NMR (CDCl₃): δ 15.00, 15.13 (SCH₂CH₃), 24.63 (NCH₂CH₂CH₂), 26.28, 26.59, 27.22 (NCH₂CH₂CH₂), 34.13 (CH₂CH₂O), 50.19 (NCH₂), 52.80 (NCHCH), 56.60 (OCH₃), 61.08 (NCH), 65.13 (CH₂CH₂O), 65.64 (OCH₂CHCH₂), 108.70 (arom. CH), 109.47 (arom. CH), 118.55 (OCH₂CHCH₂), 128.58 (CCON), 131.73 (OCH₂CHCH₂), 137.17 (CNO₂), 147.98 (CH₂CH₂OC), 154.57 (COCH₃), 166.61 (CON), 170.14 (COO). IR (Nujol) ν = 3550-2720, 3000, 2630, 2200, 1740, 1640, 1580, 1530, 1340, 1280, 1220, 1180, 1050 cm⁻¹. MS (EI): m/e (relative intensity): 527 (M⁺, 1), 377 (10), 310 (12), 309 (72), 308 (94), 268 (20), 142 (4). HRMS calcd. for C₂₄H₃₅O₇N₂S₂ = 527.1875, found = 527.1885.

5-Amino-3-(4-(2-diethylthiomethyl-(2S)-perhydro-1-pyrroloylcarbonyl)-2-methoxyphenyloxy)2-propenylpropanoate (9)

A solution of 8 (7.21 g, 14.05 mmol) and in(II) chloride (15.85 g, 76 mmol) was refluxed for 40 minutes in ethyl acetate (100 mL) then allowed to cool. The solvent was then removed *in vacuo* and the residue was triturated with saturated bicarbonate solution at 0°C. EtOAc (50 mL) was added and the reaction stirred overnight. The reaction mixture was then filtered through Celite and the filter cake washed with ethyl acetate. The combined organics were then washed with water and brine, dried with sodium sulphate and the solvent removed *in vacuo*. The product was purified using flash column chromatography (5% MeOH/dichloromethane) to give a yellow oil, (5.87g, 86%). ¹H NMR (270 MHz, CDCl₃): δ 6.82 (s, 1H, arom. CH), 6.28 (s, 1H, arom. CH), 5.99-5.85 (m, 1H, OCH₂CHCH₂), 5.31 (dd, 1H, J = 1.28 Hz, 27.66 Hz, OCH₂CHCH₂), 5.26 (dd, 1H, J = 1.28 Hz, 20.70 Hz, OCH₂CHCH₂), 4.71-4.62 (m, 5H, including doublet at 4.62, 2H, J = 5.49 Hz, NH₂ + NCHCH, OCH₂CHCH₂), 4.27 (t, 2H, J = 6.59 Hz, CH₂CH₂O), 3.92, (m, 1H, NCH), 3.74 (s, 3H, OCH₃), 3.66-3.57 (m, 2H, NCH₂) 2.89 (t, 2H, J = 6.6 Hz, CH₂CH₂O), 2.83-2.64 (m, 4H, SCH₂CH₃), 2.28-1.80 (m, 4H,

NCH₂CH₂CH₂), 1.25 (m, 6H, SCH₂CH₃); ¹³C NMR (CDCl₃) δ 14.20 (SCH₂CH₃), 26.55, 27.23 (NCH₂CH₂CH₂), 34.27 (CH₂CH₂O), 53.20 (NCHCH), 56.08 (OCH₃), 60.10 (NCH), 60.39 (NCH₂), 64.20 (CH₂CH₂O), 64.41 (OCH₂CHCH₂), 102.26 (arom. CH), 113.71 (arom. CH), 118.40 (OCH₂CHCH₂), 131.93 (OCH₂CHCH₂), 141.03 (CNH₂), 141.74 (CH₂CH₂OC), 154.56 (COCH₃), 169.69 (CON), 170.53 (COO). IR (neat liquid film) 3500-3000, 3460, 3400, 2970, 1740, 1650, 1535, 1470, 1345, 1290, 1225, 1190 cm⁻¹; MS (EI): m/e (relative intensity): 482 (M⁺, 4), 347 (2), 278 (31), 137 (1), 70 (3); HRMS calcd. for C₂₃H₃₄O₅N₂S₂ = 482.1909, found = 482.1925.

3-(4-(2-Diethylthiomethyl-(2S)-perhydro-1-pyrrolylcarbonyl)-2-methoxy-5-(2,2,2-trichloroethyloxycarbonylamino)phenyloxy)2-propenylpropanoate (10)

To a solution of **9** (5.67g, 11.74 mmol) in dichloromethane (200 mL) was added pyridine (2.02 mL, 23.48 mmol). To this was added dropwise at 0°C a solution of trichloroethyl chloroformate (1.616 mL, 11.74 mmol). The solution was stirred for a further 1 hour at 0°C. The organics were washed with 1 N HCl (3 x 100 mL), water (3 x 100 mL) brine (100 mL), dried over magnesium sulphate and the solvent removed in vacuo to give a brown oil (6.8g, 88%) ¹H NMR (270 MHz, CDCl₃): δ 9.14 (bs, 1H, NH), 7.88 (bs, 1H, CHCNH), 6.93 (s, 1H, MeOCCHC), 5.99-5.86 (m, 1H, OCH₂CHCH₂), 5.31 (dt, 1H, J = 1.47 Hz, 27.84 Hz OCH₂CHCH₂), 5.25 (dt, 1H, J = 1.29 Hz, 21.61 Hz, CH₂CHCH₂), 4.89-4.77 (m, 4H, including doublet 1H, J = 1.28 Hz, CHCHSEt, NH, CH₂-TrOC), 4.62 (d, 2H, J = 1.28 Hz, OCH₂CHCH₂), 3.81 (s, 3H, OCH₃), 3.60 (m, 2H, NCH₂), 2.91 (d, 2H, J = 6.42 Hz, CH₂CH₂O), 2.84-2.61 (m, 4H, SCH₂CH₃), 1.37-1.23 (m, 6H, SCH₂CH₃); ¹³C NMR (CDCl₃): δ 170.33 (ester CO), 168.50 (CON), 151.94 (OCO), 150.29 (COCH₃), 144.52 (COCH₂CH₂), 131.93 (OCH₂CHCH₂), 131.35 (CNH), 118.29 (OCH₂CHCH₂), 112.21 (arom. CH), 105.51 (arom. CH), 95.27 (CCl₃), 76.24 (CH₂TrOC), 74.39 (CH₂TrOC), 65.42 (CH₂CH₂O), 61.14 (NCH), 56.30 (OCH₃), 53.00 (NCHCHSEt), 34.27 (CH₂CH₂O), 27.30, 26.71, 26.43, 25.24 (NCH₂CH₂CH₂), 15.27, 14.87, 14.18 (SCH₂CH₃). MS (EI): m/e (relative intensity): 658, 656 (M⁺, 1), 508 (1), 373 (6), 305

(5), 304 (27), 192 (5), 70 (12).

3-(11-Hydroxy-5-oxo-10-(2,2,2-trichloroethyloxocarbonylamino)-
(11aS)-2,3,5,10,11,11a-hexahydro-1H-benzo[e]pyrrolo[2,1-
a][1,4]diazepin-8-yloxy-2-propenylpropanoate (6d)

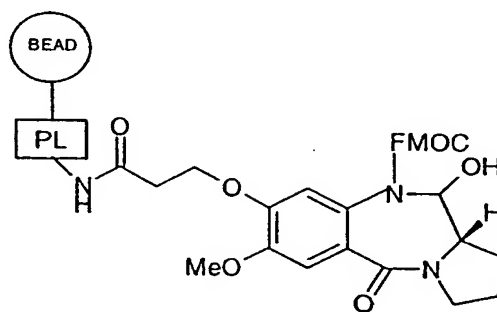
A solution of 10 (6.8g, 10.34 mmol) in acetonitrile/water (4:1, 200 mL) was treated with calcium carbonate (2.585g, 25.85 mmol) and mercuric(II) chloride (7.00g, 25.85 mmol) and the solution was stirred for 18 hours. The reaction was then filtered through Celite and the filter pad washed with ethyl acetate. The organics were collected and washed with water (3 x 50 mL), brine (100 mL) and dried over magnesium sulphate. The solvent was removed *in vacuo* and the resulting product was purified by flash column chromatography (ethyl acetate) to give the product as a yellow oil (3.67 g, 64%) ¹H NMR (270 MHz, CDCl₃): δ 7.25 (arom. CH), 6.86 (s, 1H, arom. CH), 6.00-5.85 (m, 1H, CH₂CHCH₂), 5.67 (d, 1H, J = 9.71 Hz, TrOC-CH₂) 5.37-5.20 (m, 3H, TrOC-CH₂ + OCH₂CHCH₂), 4.65 (d, 2H, J = 5.67 Hz, CH₂CHCH₂O), 4.36-4.22 (m, 3H, CH₂CH₂O + NCHOH), 3.90 (s, 3H, OCH₃), 3.72-3.47 (m, 3H, NCH + NCH₂), 2.91 (t, J = 6.41 Hz, CH₂CH₂O) 2.29-2.00 (m, 4H, NCH₂CH₂CH₂) ¹³C NMR (67.8 MHz, CDCl₃): δ 170.33 (ester carbonyl CO), 166.17 (CON), 154.4 (OCO), 149.88 (COCH₃), 148.93 (COCH₂CH₂), 131.86 (CH₂CHCH₂), 127.48 (arom. CN), 126.24 (CCON), 118.42 (OCH₂CHCH₂), 114.48 (arom. CH), 110.82 (arom. CH), 95.09 (CCl₃), 86.42 (NCHOH), 74.96 (TrOC-CH₂), 65.47 (OCH₂CHCH₂), 64.43 (CH₂CH₂O), 60.13 (NCH), 56.14 (OCH₃), 46.44 (NCH₂), 34.26 (CH₂CH₂O), 28.64 (NCH₂CH₂CH₂), MS (EI) m/z (relative intensity): = 552 (M⁺ 10), 550 (10), 374 (2), 368 (5), 304 (15), 192 (8), 70 (24), 55(24). HRMS calcd. for C₂₂H₂₃N₂O₈Cl₃ = 552.0651, found 3 peaks due to chlorine 552.0646, 550.676, 554.0617.

3-(11-Hydroxy-5-oxo-7-methoxy-10-(2,2,2-trichloroethyloxocarbonylamino)-(11aS)-2,3,5,10,11,11a-hexahydro-1H-benzo[e]pyrrolo[2,1-a][1,4]diazepin-8-yloxypropanoic acid (7d).

A solution of 6d (3.5 g, 6.35 mmol) was dissolved in ethanol

(100 mL). To this was added Tetrakis (triphenylphosphine) palladium(0) (350 mg, 0.303 mmol) and the solution refluxed for 30 minutes until the reaction was complete by TLC monitoring. The reaction was then allowed to cool and the filtered through Celite. The EtOH was then removed in vacuo to give the crude material as a yellow solid which was used directly in the next steps. ¹H-NMR (220 MHz, CDCl₃): δ 7.22 (s, 1H, OCCHCN), 7.01 (s, 1H, MeOCCHC), 6.27 (bs, COOH), 5.67 (d, 1H, J = 9.5 Hz, TrOC-CH₂), 5.06 (d, 1H, J = 12.09 Hz, TrOC-CH₂), 4.29-4.11 (m, 2H, CHOH), 3.85 (s, 3H, OCH₃), 3.71 (t, 2H, J = 6.97 Hz, CH₂CH₂O), 3.51 (m, 1H, NCH), 2.80 (m, 2H, NCH₂), 2.12-1.99 (m, 4H, NCH₂CH₂CH₂), 1.21 (t, 2H, J = 6.96 Hz, CH₂CH₂O); ¹³C NMR (67.8 MHz, CDCl₃): δ = 174.27 (acid CH), 167.34 (CON), 154.20 (OCO), 149.78 (COCH₃), 148.74 (COCH₂CH₂), 133.79 (arom. CH), 132.16 (arom. CH), 128.66 (arom. CN), 125.87 (CCON), 95.06 (CCl₃), 86.53 (NCHCHOH), 74.95 (CH₂-TrOC), 60.67 (NCH), 58.24 (CH₂CH₂O), 56.04 (OCH₃), 46.44 (NCH₂), 35.24 (NCH₂CH₂CH₂), 28.59 (NCH₂CH₂CH₂), 23.08 (CH₂CH₂O).

Example 1(e): Synthesis of resin-bound protected PBD (JGB-285)

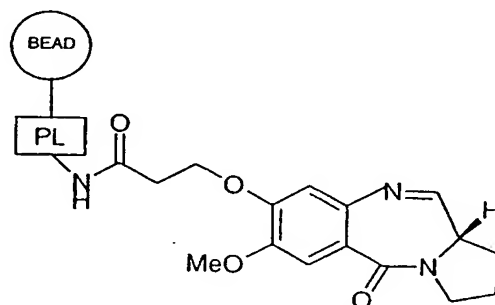


JGB-285

DMF (500 µl) was added to amino Tentagel resin (0.056 g, 0.26 mmol/g loading) in an Alltech tube (8 ml) and the resulting suspension shaken for 30 min. A solution of Fmoc-PBD-acid **7b** (66.6 mg, 0.12 mmol), TBTU (0.038 g, 0.012 mmol) and DIPEA (21 µl, 0.012 mmol) in DMF (1 ml) was added and shaking continued for 16 hours. The resin was filtered and rinsed with DMF (5 ml), CH₂Cl₂ (5 ml) and MeOH (5 ml). This procedure was repeated twice to ensure complete reaction and the resin

was then dried *in vacuo* to afford JGB-285.

Example 1(f): Synthesis of resin-bound unprotected PBD (JGB-286)



JGB-286

A solution of piperidine in DMF (20%, 500 μ l) was added to the resin JGB-285 and the suspension shaken for 16 hours. The resin was filtered and rinsed with DMF (5 ml), CH_2Cl_2 (5 ml) and MeOH (5 ml). This procedure was repeated twice and the resin was dried *in vacuo* to afford JGB-286.

Example 2: Synthesis of 8-aminopropyl PBD of formula I (see Figure 2)

Overall Synthesis

The compound 19 was prepared by removal of Fmoc from 18 under standard conditions (piperidine/DMF). The Fmoc carbamate was obtained via Swern oxidation of the alcohol 17, which resulted in spontaneous closure of the pyrrolobenzodiazepine B-ring. A number of other oxidation methods should also prove effective in promoting the oxidation/cyclization reaction, for example, the Dess Martin reagent, the TPAP/NMO system or Pyridine Sulphur trioxide in DMSO. The alcohol 17 was furnished by treatment of the amino alcohol 16 with Nvoc-Cl in the presence of pyridine. As before this is a general procedure applicable to any chloroformate, the choice limited only by compatibility with the PBD and Fmoc cleavage conditions. The amine group can also be protected with numerous other carbamate protecting groups, the most useful in this instance being Alloc, Teoc and Noc due to their compatibility with Fmoc cleavage conditions.

It should be noted that Fmoc itself could be employed for N-10 protection in which case it would obviously be necessary to employ a different protecting group for the aliphatic nitrogen (see below). The amino alcohol was prepared by tin chloride reduction of the nitro alcohol, which in turn was prepared by coupling pyrrolidine methanol to the o-nitrobenzoic acid **14** under standard conditions. The o-nitrobenzoic acid was prepared by Fmoc protection of the amino acid **13**. Again, it would be possible to substitute Fmoc with a number of other protecting groups for example Boc, Alloc, Noc etc. due to their compatibility with the N10 Nvoc group. It should be noted that if Fmoc was used to protect the aromatic N10 group, Boc, Alloc, Teoc and Nvoc could be used to protect the aliphatic nitrogen. The amino acid **13** was prepared by hydrolysis of the ester **12**, which in turn was obtained by simultaneous nitration and deprotection of the Boc protected amine **11**, which was obtained by a Mitsunobu etherification of methyl vanillate with Boc aminopropanol.

Boc Amino Ester (11)

A solution diethylazidodicarboxylate (3.38 g, 19.4 mmol) in THF (50 ml) was added dropwise to a solution of methylvanillate (3.53 g, 19.4 mmol), N-Boc-propanolamine (3.4 g, 19.4 mmol) and triphenylphosphine (5.09 g, 19.4 mmol) in THF (50 ml) at 0°C. The reaction mixture was allowed to warm to room temperature and stir overnight. Excess solvent was removed by rotary evaporation under reduced pressure and the residue triturated with toluene. Precipitated triphenylphosphine oxide was removed by vacuum filtration and the filtrate concentrated *in vacuo*. The residue was subjected to flash column chromatography (silica gel, petroleum ether 40 - 60/ethyl acetate, 80/20) and removal of excess eluent afforded the pure product **11** (4.8 g, 73 % yield.). ¹H NMR (270 MHz, CDCl₃) δ 7.65 (dd, J = 8.43, 2.02 Hz, 1H), 7.54 (d, J = 2.02 Hz, 1H), 6.86 (d, J = 8.43 Hz, 1H), 5.55 (bs, 1H), 4.15 (t, J = 5.87 Hz, 2H), 3.93 (s, 3H), 3.90 (s, 3H), 3.41-3.35 (m, 2H), 2.09-2.00 (m, 2H) and 1.46 (s, 9H). ¹³C NMR (68.7 MHz, CDCl₃) δ 166.9, 156.1, 152.1, 148.8, 123.5, 122.8, 112.0,

111.2, 79.0, 68.2, 55.9, 52.0, 38.9, 29.2 and 28.5.

Amino Nitro Ester (12)

5 The Boc-protected amine 11 (10 g) was added portionwise to cold nitric acid (30 ml, 70%, ice bath), the reaction mixture was allowed warm to room temperature and stir overnight. The reaction mixture was poured onto crushed ice (100 g) and the resulting aqueous solution reduced to half its original volume by rotary evaporation under reduced pressure. The resulting
10 precipitate was collected by vacuum filtration and recrystallised from absolute ethanol to afford the product as a yellow crystalline solid 12 (8.9 g, 87 %). ¹H NMR (270 MHz, CDCl₃) δ 7.47 (s, 1H), 7.08 (s, 1H), 4.24 (t, J = 5.86 Hz, 2H), 3.96, (s, 3H), 3.89 (s, 3H), 3.24 (t, J = 6.78 Hz, 2H) and
15 2.32-2.23 (m, 2H).

Amino Nitro Acid (13)

A solution of potassium hydroxide (0.5 g, 8.7 mmol) and the nitrobenzoic acid 12 (1 g, 2.9 mmol) in aqueous methanol (H₂O, 10 ml; methanol, 20 ml) was allowed to stir at room
20 temperature for 1 hour and then heated at reflux until TLC (AcOEt, MeOH, TEA, 1:10:100) revealed the complete consumption of starting material. Excess methanol was removed by rotary evaporation and the residual solution diluted with water and
25 neutralised with 1N HCl. The neutralised aqueous solution was used directly, without further purification, in the next synthetic step.

Fmoc Nitro Acid (14)

30 Fluorenylmethyl chloroformate (0.78 g, 3 mmol) was added portionwise to the aqueous solution from the previous reaction which had been diluted with THF (50 ml) and aqueous sodium carbonate (2.15 g, 50 ml water). The reaction mixture was then allowed to stir overnight. Excess organic solvent was
35 removed by rotary evaporation under reduced pressure from the reaction mixture, the residual aqueous solution was then washed with ethyl acetate (3 x 20 ml) (to remove excess Fmoc-Cl). The aqueous phase was acidified with conc. HCl and

extracted with ethyl acetate (2 x 50 ml). The organic phase was dried over magnesium sulphate, filtered and evaporated in vacuo to afford the product **14** (1 g, 70% yield). ¹H NMR (270 MHz, CDCl₃) δ (Rotamers) 8.21 (bs, 2H), 7.73 (d, J = 7.14 Hz, 2H), 7.59 (d, J = 7.33 Hz, 2H) 7.40 - 7.13 (m, 5H), 6.47 and 5.70 (2 x bs, 1H), 4.54-3.88 (m, 5H), 3.77 (s, 3H), 3.44-3.42 (m, 2H) and 2.04-1.90 (m, 2H). ¹³C NMR (68.7 MHz, CDCl₃) δ 168.7, 156.9, 152.1, 149.8, 143.7, 141.9, 141.3, 127.7, 127.0, 124.9, 120.6, 120.0, 111.1, 107.8, 68.5, 66.4, 56.4, 47.3, 39.1 and 28.4.

Fmoc Nitro Alcohol (**15**)

A catalytic amount of DMF (2 drops) was added to a solution of the acid **14** (1.16 g, 2.36 mmol) and oxalyl chloride (0.33 g, 2.6 mmol) in dry dichloromethane (20 ml) and the reaction mixture was allowed to stir overnight. The resulting acid chloride solution was cooled to 0°C and treated dropwise with a solution of pyrrolidinemethanol (0.26 g, 2.57 mmol) and triethylamine (0.52 g, 5.14 mmol) in dry dichloromethane (15 ml). Thin layer chromatography, performed shortly after the end of the addition of amine, revealed that reaction had gone to completion. The reaction mixture was washed with HCl (1N, 1 x 50 ml) and water (2 x 20 ml) and dried over magnesium sulphate. Removal of excess solvent afforded the crude product which was subjected to flash column chromatography (silica gel, gradient elution, 1% methanol in chloroform to 2% methanol in chloroform) to afford the required amide **15** (1.1 g, 81%). ¹H NMR (270 MHz, CDCl₃) δ 7.75 (d, J = 7.33 Hz, 2H), 7.67 (s, 1H), 7.60 (d, J = 6.96 Hz, 2H), 7.41-7.26 (m, 4H), 6.78 (s, 1H), 5.66 (bs, 1H), 4.48-4.39 (m, 3H), 4.23-4.13 (m, 3H), 3.91-3.79 (m, 5H), 3.45-3.42 (m, 2H), 3.18-3.13 (m, 2H) and 2.08-1.70 (m, 6H). ¹³C NMR (68.7 MHz, CDCl₃) δ 168.5, 156.5, 154.7, 148.2, 143.9, 141.3, 137.0, 128.0, 127.7, 127.0, 124.9, 120, 108.9, 108.0, 68.4, 66.2, 66.0, 61.5, 56.6, 53.5, 47.3, 39.0, 28.9, 28.4 and 24.4.

Fmoc Amino Alcohol (16)

A solution of the nitroamide 15 (3 g, 5.22 mmol) and $\text{SnCl}_4 \cdot 2\text{H}_2\text{O}$ (6.15 g, 27.15 mmol) in methanol (60 ml) was heated at reflux for 2 hours. The reaction mixture was concentrated to 1/3 of its original volume and carefully treated with saturated aqueous sodium bicarbonate solution (vigorous effervescence!) until pH8 was obtained. The mixture was allowed to stir vigorously with ethyl acetate (100 ml) overnight and then filtered through celite to remove precipitated tin salts. The aqueous phase was extracted with ethyl acetate (50 ml) and the combined organic phase was dried over magnesium sulphate. Removal of excess solvent afforded the desired amine as a dark yellow oil 16 (1.93 g, 68%). ^1H NMR (270 MHz, CDCl_3) δ 7.75 (d, J = 7.51 Hz, 2H), 7.61 (d, J = 7.33 Hz, 2H), 7.40-7.26 (m, 4H), 6.72 (s, 1H), 6.25 (s, 1H), 5.95 (bs, 1H), 4.43-4.04 (m, 6H), 3.67-3.42 (m, 9H) and 2.11-1.7 (m, 6H). ^{13}C NMR (68.7 MHz, CDCl_3) δ 171.7, 156.6, 150.8, 144.0, 141.3, 140.6, 127.6, 127.0, 125.0, 119.9, 112.0, 102.2, 68.0, 66.6, 66.4, 61.0, 56.6, 51.0, 47.3, 39.5, 29.1, 28.5 and 24.9.

Fmoc Nvoc Alcohol (17)

A solution of 4,5-dimethoxy-2-nitrobenzylchloroformate (1.44 g, 5.23 mmol) in dichloromethane (40 ml) was added dropwise to a solution of the amine 16 (2.59 g, 4.75 mmol) and pyridine (0.41 g, 5.23 mmol) in dichloromethane (60 ml) at 0°C. After 3 hours the reaction mixture was washed with HCl (1N, 2 x 100 ml), water (2 x 100 ml) and brine (1 x 100 ml). The organic phase was dried over magnesium sulphate and removal of excess solvent gave the crude product, which was subjected to flash column chromatography (silica gel, ethyl acetate followed by 1% methanol in ethyl acetate) to afford the pure carbamate 17 (3.2 g, 86%). ^1H NMR (270 MHz, CDCl_3) δ 8.94 (br, s 1H) 7.74 (d, J = 7.51 Hz, 2H), 7.71 (s, 1H), 7.61 (d J = 7.33 Hz, 2H), 7.40-7.25 (m, 4H), 7.08 (s, 1H), 6.80 (s, 1H), 5.62 (d, J = 15.02 Hz, 1H), 5.50 (d, J = 15.02, 1H), 4.44-4.41 (m, 3H), 4.24-4.13 (m, 3H), 3.99 (s, 3H), 3.94 (s, 3H), 3.70-3.44 (m, 9H), and 2.17-1.72 (m, 6H). ^{13}C NMR (68.7 MHz, CDCl_3) δ 171.7, 164.0, 156.6, 153.7, 153.3, 150.1, 148.1, 144.3, 144.0, 141.3,

139.6, 131.3, 127.6, 127.0, 125.0, 119.9, 110.7, 110.1, 108.2, 105.5, 68.1, 66.4, 66.1, 63.9, 60.9, 56.6, 56.4, 56.2, 47.3, 39.5, 28.9, 28.3 and 25.1.

5 **Fmoc Nvoc Carbinolamine (18)**

A solution of DMSO (0.8 ml, 11.4 mmol) in dry dichloromethane (15 ml) was added dropwise, over 30 minutes, to a solution of oxalyl chloride (0.72 g, 5.71 mmol) in dry dichloromethane (15 ml) at -45°C under a nitrogen atmosphere. The reaction mixture was allowed to stir for 30 minutes before the addition of the substrate **17** (3.2 g, 4.08 mmol) in dry dichloromethane (35 ml) over 50 minutes whilst maintaining the temperature of the reaction at -45°C. The reaction mixture was then allowed to stir at -45°C for a further 45 minutes. A solution of triethylamine (2.15 ml, 16.2 mmol) in dry dichloromethane (10 ml) was added dropwise over 25 minutes at -45°C and the reaction mixture allowed to stir for a further 30 minutes at -45°C before being allowed to warm to room temperature. The reaction mixture was washed with 1N HCl (1 x 75ml), water (1 x 75 ml), brine (1 x 75 ml) and dried over magnesium sulphate. Removal of excess solvent furnished the crude product which was subjected to flash column chromatography (silica gel, ethyl acetate) to afford the cyclized product **18** (1.92 g, 60% yield). ¹H NMR (270 MHz, CDCl₃) δ 7.74 (d, J = 7.51 Hz, 2H), 7.60-7.59 (m, 3H), 7.40-7.23 (m, 4H), 7.22 (s, 1H), 6.83 (s, 1H), 6.50 (s, 1H), 5.88 (bs, 1H), 5.72 (d, J = 9.34 Hz, 1H), 5.45-5.38 (m, 2H), 4.59 (bs, 1H), 4.42 (d, J = 7.14 Hz, 2H), 4.22-4.08 (m, 3H), 3.86 (s, 3H), 3.76 (s, 3H), 3.68 (s, 3H), 3.59-3.44 (m, 4H) and 2.12-2.02 (m, 6H). ¹³C NMR (68.7 MHz, CDCl₃) δ 166.9, 156.6, 155.4, 153.8, 150.1, 148.8, 148.1, 144.0, 141.3, 139, 128.2, 127.7, 127.0, 126.7, 126.5, 125.0, 120.0, 113.7, 110.5, 109.7, 108.1, 86.2, 68.4, 66.3, 65.4, 60.3, 56.3, 56.2, 56.0, 47.3, 46.5, 39.4, 29.7, 28.7 and 23.1.

35 **Amino Nvoc Carbinolamine (19)**

The Fmoc protected amine **18** (0.5 g, 0.64 mmol) was added to a solution of piperidine (1 g, 11.7 mmol) in dichloromethane (10 ml). After 2 hours TLC revealed the continued presence of

starting material DMF was added as a co-solvent and reaction proceeded to completion over the next 30 minutes. The reaction mixture was diluted with ethyl acetate (50 ml), washed with water (25 ml), brine (25 ml) and dried over magnesium sulphate. Removal of excess solvent afforded a yellow crystalline product which was recrystallised from ethyl acetate and petroleum ether 40-60 (0.123 g, 34% yield). ¹H NMR (270 MHz, CDCl₃) δ 7.61 (s, 1H), 7.21 (s, 1H), 6.90 (s, 1H), 6.50 (s, 1H), 5.70 (d, J = 9.70 Hz, 1H), 5.44 (bs, 2H), 4.13-4.11 (m, 1H), 3.96-3.40 (m, 13H), and 2.18-1.90 (m, 6H). ¹³C NMR (68.7 MHz, CDCl₃) δ 167.1, 155.1, 153.8, 150.0, 148.7, 147.9, 138.8, 128.5, 127.3, 126.5, 114.3, 110.4, 109.5, 107.9, 86.0, 67.1, 65.1, 60.7, 56.3, 56.1, 53.3, 38.7, 28.7, 28.0 and 23.1.

Example 3a: Synthesis of PBD-Triglycine 30 (Figures 3a & 3b)

This example was carried out to prove the general method of synthesis.

Resin Deprotection

Fmoc-aminoethyl photolinker NovaSyn TG resin 20 (0.35 g, 0.23 mmol/g loading) was placed in a peptide vessel, fitted with a sinter. After the addition of 20% piperidine in DMF (3 ml), the vessel was shaken for 3 hours. The deprotected resin 21 was then separated by filtration and rinsed with NMP (3 ml), MeOH (3 ml) and CH₂Cl₂ (3 ml). The whole procedure was repeated twice before drying the resin *in vacuo*.

Coupling conditions

DMF (2 ml) was added to resin 21 and the suspension shaken for 30 min. A solution of Fmoc-glycine (0.24 g, 0.805 mmol), TBTU (0.26 g, 0.805 mmol) and DIPEA (140 μl, 0.805 mmol) in DMF (2ml) was added and shaking continued for 20 hours. The coupled resin 22 was then filtered and rinsed with NMP (5 ml), MeOH (5 ml) and CH₂Cl₂ (5 ml). The whole procedure was repeated once before drying the resin *in vacuo*. The coupling efficiency was monitored by the addition of bromophenol blue in DMF (0.2 ml).

Acetylation (Endcapping) conditions

Ac₂O (20%) and pyridine (30%) in CH₂Cl₂ (5 ml) was added to resin 22 and the suspension was shaken for 2 hours. The acetylated resin was filtered and washed with CH₂Cl₂ (5 ml),
5 EtOH (5 ml) and a further aliquot of CH₂Cl₂ (2 ml). The whole procedure was repeated once before drying the resin *in vacuo*. The effectiveness of acetylation was monitored by the addition of bromophenol blue in DMF (0.5 ml).

10 Deprotection conditions

Piperidine (20%) in DMF (2 ml) was added to the acetylated resin 22 and the suspension was shaken for 12 hours. The deprotected resin 23 was collected by filtration and rinsed with NMP (5 ml), MeOH (5 ml) and CH₂Cl₂ (5 ml). The whole
15 procedure was repeated twice before drying the resin *in vacuo*.

Addition of Two Further Glycine Units

The previous coupling (21 - 22), acetylation and deprotection (22 - 23) steps were repeated twice (23 - 27) until the resin-bound tripeptide 27 was obtained.
20

Coupling to the PBD Unit (Fmoc-PBD)

Triglycine resin 27 (0.12 g, 0.235 mmol/g loading) was placed in a peptide synthesis vessel fitted with a sinter. DMF (3
25 ml) was added and the vessel was shaken for 30 min. A solution of the Fmoc-PBD acid 7b (0.15 g, 0.28 mmol), TBTU (0.09 g, 0.28 mmol) and DIPEA (50 µl, 0.28 mmol) in DMF (3 ml) was added, and shaking continued for 20 hours. Bromophenol blue indicator (30 µl) was added to monitor the progress of
30 the reaction. The coupled resin 28b was collected by filtration and rinsed with DMF (5 ml), NMP (5 ml) and CH₂Cl₂ (5 ml). The whole procedure was repeated twice before drying the resin *in vacuo*.

35 Acetylation (Endcapping) Conditions

Ac₂O (20%) and pyridine (30%) in CH₂Cl₂ (5 ml) was added to resin 28b and the vessel shaken for 2 hours. The acetylated resin was filtered and washed with CH₂Cl₂ (5 ml), EtOH (5 ml)

and further CH_2Cl_2 (2 ml). The whole procedure was repeated once and the resin 29 dried *in vacuo*. The effectiveness of acetylation was monitored by the addition of bromophenol blue in DMF (0.5 ml).

5

Deprotection to Free PBD

Piperidine (20%) in DMF (2 ml) was added to the acetylated resin 28a and the vessel shaken for 12 hours. The resin 29 was collected by filtration and rinsed with NMP (5 ml), MeOH (5 ml) and CH_2Cl_2 (5 ml). This procedure was repeated twice and the resin dried *in vacuo*.

10

Synthesis of Nvoc-PBD Triglycine 28a (Figure 3b)

Triglycine resin 27 (0.16 g, 0.235 mmol/g loading) was placed in a peptide vessel fitted with a sinter. DMF (3 ml) was added and the vessel shaken for 30 min. A solution of the Nvoc-PBD acid 7a (0.22 g, 0.38 mmol), TBTU (0.12 g, 0.38 mmol) and DIPEA (65 μl , 0.38 mmol) in DMF (3 ml) was added, and shaking continued for 20 hours. Bromophenol blue indicator (30 μl) was added to monitor the progress of the reaction. Resin 28a was collected by filtration and rinsed with DMF (5 ml), NMP (5 ml) and CH_2Cl_2 (5 ml). The whole procedure was repeated twice before drying the resin *in vacuo*.

15

20

Acetylation (Endcapping) Conditions

Ac_2O (20%) and pyridine (30%) in CH_2Cl_2 (5 ml) was added to resin 28a and the vessel shaken for 2 hours. The acetylated resin was collected by filtration and washed with CH_2Cl_2 (5 ml), EtOH (5 ml) and further CH_2Cl_2 (2 ml). The whole procedure was repeated once before drying the resin 29 *in vacuo*. The effectiveness of acetylation was monitored by the addition of bromophenol blue in DMF (.5 ml).

30

Photolysis

A suspension of beads bearing the Nvoc-PBD in DMF was simultaneously N10-deprotected and cleaved from the resin by irradiating at 365 nm for 2 hours (Spectrolinker XL 1000 UV

35

Crosslinker, Spectronics Corporation) to afford a 1 mmol stock solution of 30 which was used directly in the MTT assay (Example 3b).

5 Example 3(b): General MTT Assay Method

The ability of agents to inhibit the growth of U937 chronic human histiocytic leukemia cells or K562 human chronic myeloid leukemia cells in culture was measured using the MTT assay (Mosmann, 1983). This is based on the ability of viable cells to reduce a yellow soluble tetrazolium salt, 3-(4,5-dimethylthiazolyl)-2,5-diphenyltetrazolium bromide (MTT; Sigma Chemical Co.), to an insoluble purple formazan precipitate. Following drug treatment, the cells were transferred to 96-well microtitre plates with 10^4 cells per well and 8 wells per sample. The plates were incubated at 37°C in a humidified atmosphere containing 5% CO₂. Following incubation of the plates for 4 days (to allow control cells to increase in number by 10-fold), 20 µL of a 5mg/mL solution of MTT in phosphate-buffered saline was added to each well and the plates incubated further for 5 hours. The plates were then centrifuged for 5 minutes at 300g and the bulk of the medium removed from the cell pellet, leaving 10-20 µL per well. DMSO (200 µL) was added to each well, and the samples agitated to ensure complete mixing. The optical density was then read at a wavelength of 550nm on a Titertek Multiscan ELISA plate reader and the dose-response curve constructed. The IC₅₀ value was read as the dose required to reduce the final optical density to 50% of the control value.

30 MTT Assay of PBD-Triglycine

The MTT assay was used to evaluate the cytotoxicity of the previously prepared PBD-Triglycine (compound 30), from Example 3a. The assay found the IC₅₀ to be 0.59 µM.

35 Example 4(a): Synthesis of the Resin-bound Tripeptide Library (Figures 4a and 4b)

Following Example 3, this procedure was used to synthesize a tripeptide library.

Resin Deprotection

Fmoc-aminoethyl photolinker NovaSyn TG resin **20** (1.35 g, 0.23 mmol/g loading) was weighed into 27 Alltech tubes in 50 mg portions. Piperidine (20%) in DMF (0.5 ml) was added to each tube, and the tubes were then placed onto an orbital shaker for 3 hours. The deprotected resin **21** was collected by filtration using a Supelco Vacuum Manifold and rinsed with DMF (2 ml) and CH₂Cl₂ (2 ml). The whole procedure was repeated twice before drying the resin *in vacuo*.

Coupling conditions

DMF (0.5 ml) was added to each Alltech tube containing resin **21**, and the tubes were shaken for 30 min. Fmoc-glycine (0.306 g, 1.03 mmol) in DMF (3.4 ml) was added to the first 9 tubes; Fmoc-valine (0.36 g, 1.06 mmol) in DMF (3.54 ml) to the next 9 tubes and Fmoc-phenylalanine (0.396 g, 1.03 mmol) in DMF (3.4 ml) to the final 9 tubes. TBTU (0.972 g, 3.03 mmol) in DMF (10 ml) and DIPEA (540 µl, 3.1 mmol) were added to all 27 tubes and shaking was continued for 20 hours. The coupled resin **31** was collected by filtration and rinsed with DMF (5 ml), CH₂Cl₂ (5 ml) and EtOH (5 ml). The whole procedure was repeated twice before drying the resin *in vacuo*. The coupling efficiency was monitored by the addition of a few drops of 10% DIPEA/DMF and 1% TNBS/DMF. The resin remained colourless when coupling was complete.

Acetylation (Endcapping) Conditions

Ac₂O (20%) and pyridine (30%) in CH₂Cl₂ (1 ml) were added to each Alltech tube containing resin **31**, and the tubes were shaken for 2 hours. The acetylated resin was filtered and washed with CH₂Cl₂ (5 ml), EtOH (5 ml) and further CH₂Cl₂ (2 ml). The whole procedure was repeated once before drying the resin *in vacuo*.

Deprotection Conditions

Piperidine (20%) in DMF (0.5 ml) was added to each tube of acetylated resin **31**, and the tubes shaken for 12 hours. The deprotected resin **32** was collected by filtration and rinsed

with DMF (5 ml) and CH_2Cl_2 (5 ml). The whole procedure was repeated twice before drying the resin *in vacuo*.

Coupling of Two Further Amino Acid Units

The previous coupling (21 - 31), acetylation and deprotection (31 - 32) steps were repeated twice using the appropriate Fmoc-protected Amino Acids in each Alltech tube (32 - 36) to achieve all possible combinations. This resulted in the resin-bound tripeptide library 36.

Coupling to the Fmoc-PBD Unit (Figure 4b)

DMF (0.5 ml) was added to each Alltech tube containing resin 36 and the tubes were shaken for 30 min. Fmoc-PBD acid 7b (0.866 g, 1.55 mmol) in DMF (5.2 ml), TBTU (0.486 g, 1.55 mmol) in DMF (5 ml) and DIPEA (270 μl , 1.55 mmol) were added and shaking was continued for 20 hours. The coupled resin 37 was collected from each tube by filtration and rinsed with CH_2Cl_2 (5 ml), EtOH (5 ml) and further CH_2Cl_2 (2 ml). The whole procedure was repeated twice before drying the batches of resin *in vacuo*.

Acetylation (Endcapping) Conditions

Ac_2O (20%) and pyridine (30%) in CH_2Cl_2 (1 ml) were added to each tube of resin 37 and the tubes were shaken for 2 hours. Acetylated resin from each Alltech tube was collected by filtration and washed with CH_2Cl_2 (5 ml), EtOH (5 ml) and further CH_2Cl_2 (2 ml). The whole procedure was repeated once before drying the resin *in vacuo*. The effectiveness of acetylation was monitored by the addition of a few drops of 10% DIPEA/DMF and 1% TNBS/DMF.

Deprotection Conditions

Piperidine (20%) in DMF (0.5 ml) was added to each tube of acetylated resin 37 and the tubes were shaken for 12 hours. The batches of deprotected resin 38 were collected by filtration and rinsed with DMF (5 ml), CH_2Cl_2 (5 ml) and MeOH (5 ml). The whole procedure was repeated twice before drying the residue *in vacuo* to afford batches of resin 38.

Photolysis

A suspension of beads bearing the deprotected resin 38 were cleaved from the resin by irradiating at 365 nm for 2 hours (Spectrolinker XL 1000 UV Crosslinker, Spectronics Corporation) to afford a 1 mmol stock solution which was used directly in the MTT assay (Example 4b).

Example 4(b): Screening of a 27 member combinatorial library prepared on beads

The combinatorial library synthesised in example 4(a) was screened using the MTT assay as previously described.

Compound	Amino-acid Sequence of Combinatorial Units	IC ₅₀ (μM)
1	GGG	0.59
2	GGV	0.63
3	GGF	0.56
4	GVG	0.55
5	GVV	0.52
6	GVF	0.64
7	GFG	0.63
8	GFV	0.78
9	GFF	0.63
10	VGG	0.59
11	VGV	0.33
12	VGF	0.58
13	VVG	0.58
14	VVV	0.58
15	VVF	0.53
16	VFG	0.46
17	VFV	0.56
18	VFF	0.56
19	FGG	0.54
20	FGV	0.54
21	FGF	0.57
22	FVG	0.58
23	FVV	0.54
24	FVF	0.69
25	FFG	0.54
26	FFV	0.40
27	FFF	0.59
GW/613	-	0.33

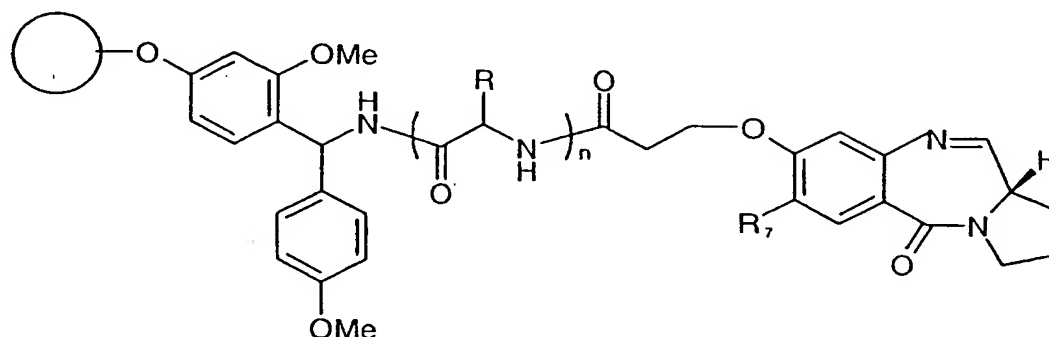
G=GLYCINE

V=VALINE

F=PHENYLALANINE

These results demonstrate that varying the amino acid sequence affects the cytotoxicity of the PBDs.

Example 5(a): Preparation of a 27-Member Tripeptide-PBD Library on Crowns for Solution Phase Testing



38: $n=3$; $R=H$, $CH_2CH(CH_3)_2$ or CH_2Ph

A 27-member library was prepared on Chiron Technology crowns from Fmoc-protected Glycine, Leucine and Phenylalanine building blocks, and an NVOC-protected PBD unit using the Multipin™ Synthesis Kit and the same general reaction scheme as exemplified in Example 4(a).

Crown Deprotection

Twenty-seven Fmoc-Rink amide O-series crowns (loading: 2.2 μM per crown) were attached to the first twenty-seven pins of a 98-pin block. The block was inverted and placed in a vessel containing a solution of piperidine (20%) in DMF (50 mL, anhydrous) on a shaker (Heidolph, Titramax 100). After 30 min, the block was removed from the container and excess piperidine/DMF allowed to drain away. The block was then inverted, placed in a vessel containing fresh DMF (50 mL), and the whole assembly agitated for 5 min. Finally, the crowns were washed twice for 2 minutes with methanol before allowing to air-dry for 20 min.

Preparation of Activated Amino Esters

A solution of N-Fmoc-Glycine (128.4 mg, 0.43 mmol), DIC (55 mg, 0.43 mmol) and HOBT (70 mg, 0.52 mmol) in DMF (2160 μ L) was

agitated for 20 min, and aliquots (200 μ L) were added to the first nine wells (H1-H9) of a deep 96-well microtitre plate. Similar solutions of active esters were prepared from N-Fmoc Leucine (152.7 mg) and N-Fmoc-Phenylalanine (167.4 mg), and aliquots (200 μ L) were dispensed into wells H10-G6 and G7-F3, respectively (see Table 1 below).

	H	G	F
1	Glycine	Leucine	Phenylalanine
2	Glycine	Leucine	Phenylalanine
3	Glycine	Leucine	Phenylalanine
4	Glycine	Leucine	
5	Glycine	Leucine	
6	Glycine	Leucine	
7	Glycine	Phenylalanine	
8	Glycine	Phenylalanine	
9	Glycine	Phenylalanine	
10	Leucine	Phenylalanine	
11	Leucine	Phenylalanine	
12	Leucine	Phenylalanine	

Table 1: Distribution of N-Fmoc-protected Amino Acids into Wells H1 to F3.

Coupling Reaction

Each well was doped with a small amount of bromophenol blue indicator (25 μ L of 6.6 mg in 10 mL of DMF) and the crown array immersed in the wells. The crowns (previously colourless) instantly turned blue indicating the presence of free amines. The block/deep well plate assembly was agitated on a shaker for 18 hours after which time all of the crowns became virtually colourless indicating that the coupling reactions had gone to completion. The crown array was then removed from the microtitre plate, excess coupling reagent allowed to drain, and the crown array washed once with DMF (50 mL, five min) and twice with methanol (100 mL, 2 min). Finally, the crown array was allowed to air dry for 20 min.

Deprotection

The crown array was inverted and placed in a vessel containing a solution of piperidine (20%) in DMF (50 mL, anhydrous) on a shaker (Heidolph, Titramax 100). After 30 min, the block was

removed from the container and excess piperidine/DMF allowed to drain away. The block was then inverted, placed in a vessel containing fresh DMF (50 mL) and the whole assembly agitated for 5 min. Finally, the crowns were washed twice for 2 minutes with methanol before allowing to air-dry for 20 min.

Coupling to the Second Amino Acid

The deprotected crown array was immersed in a deep well microtitre plate charged with freshly prepared solutions of the activated esters of N-Fmoc-Glycine, N-Fmoc-Leucine and Fmoc-Phenylalanine according to the pattern shown in Table 2 below. The block/deep well plate assembly was agitated on an orbital shaker for 18 hours after which time all of the crowns had become virtually colourless indicating that the coupling reactions had gone to completion. The crown array was then removed from the microtitre plate, excess coupling reagent allowed to drain, and the crown array washed once with DMF (50 mL, five min) and twice with methanol (100 mL, 2 min). Finally, the crown array was allowed to air dry for 20 min.

	H	G	F
1	Glycine	Leucine	Phenylalanine
2	Glycine	Leucine	Phenylalanine
3	Glycine	Leucine	Phenylalanine
4	Leucine	Phenylalanine	
5	Leucine	Phenylalanine	
6	Leucine	Phenylalanine	
7	Phenylalanine	Glycine	
8	Phenylalanine	Glycine	
9	Phenylalanine	Glycine	
10	Glycine	Leucine	
11	Glycine	Leucine	
12	Glycine	Leucine	

Table 2: Distribution of N-Fmoc-Protected Amino Acids into Wells H1 to F3.

Deprotection

The crown array was inverted and placed in a container charged with a solution of piperidine (20%) in DMF (50 mL, anhydrous) on a shaker (Heidolph, Titramax 100). After 30 min, the block was removed from the container and excess piperidine/DMF

allowed to drain away. The block was then inverted, placed in a vessel containing fresh DMF (50 mL) and the whole assembly agitated for 5 min. Finally, the crowns were washed twice for 2 minutes with methanol before allowing to air-dry for 20 min.

5

Coupling of the Third Amino Acid Unit

The deprotected crown array was immersed in a deep well microtitre plate charged with freshly prepared solutions of the activated esters of N-Fmoc-Glycine, N-Fmoc-Leucine and Fmoc-Phenylalanine according to the pattern shown in Table 3 below. The block/deep well plate assembly was agitated on an orbital shaker for 18 hours after which time all of the crowns had become virtually colourless indicating that the coupling reactions had gone to completion. The crown array was then removed from the microtitre plate, excess coupling reagent allowed to drain, and the crown array washed once with DMF (50 mL, five min) and twice with methanol (100 mL, 2 min). Finally, the crown array was allowed to air dry for 20 min.

20

25

30

	H	G	F
1	Glycine	Glycine	Glycine
2	Leucine	Leucine	Leucine
3	Phenylalanine	Phenylalanine	Phenylalanine
4	Glycine	Glycine	
5	Leucine	Leucine	
6	Phenylalanine	Phenylalanine	
7	Glycine	Glycine	
8	Leucine	Leucine	
9	Phenylalanine	Phenylalanine	
10	Glycine	Glycine	
11	Leucine	Leucine	
12	Phenylalanine	Phenylalanine	

Table 3: Distribution of N-Fmoc-protected Amino Acids into Wells H1 to F3.

35

Deprotection

40

The crown array was inverted and placed in a vessel containing a solution of piperidine (20%) in DMF (50 mL, anhydrous) on a shaker (Heidolph, Titramax 100). After 30 min, the block was removed from the container and excess piperidine/DMF allowed to drain away. The block was then inverted, placed in a vessel

containing fresh DMF (50 mL) and the whole assembly agitated for 5 min. Finally, the crowns were washed twice for 2 minutes with methanol before allowing to air-dry for 20 min.

5 Attachment of PBD Capping Unit

A solution of Nvoc-PBD acid (745 mg, 1.29 mmol), DIC (16 mg, 1.29 mmol) and HOBt (209 mg, 1.55 mmol) in DMF (6.48 ml) was agitated for 20 minutes and then dispensed into all twenty-seven wells. The block/deep well microtitre plate assembly was
10 agitated on a shaker for 48 hours. The crown array was then removed from the microtitre plate, excess coupling reagent allowed to drain, and the crown array washed once with DMF (50 mL, five min) and twice with methanol (100 mL, 2 min). Finally, the crown array was allowed to air dry for 20 min.

15 Cleavage

Prior to cleavage, the crowns were washed successively with DMF, toluene, methanol and dichloromethane to remove any non-covalent contaminants. The crown array was then immersed in
20 twenty-seven racked (but individual) 1 mL polypropylene tubes each containing TFA/H₂O (300 µL, 95:5, v/v), and the block/rack assembly was agitated on an orbital shaker for 2 hours at room temperature. Excess TFA was removed by parallel evaporation under nitrogen (supplied by a glass manifold with 8 outlets)
25 followed by final drying in vacuo over 48 hours to afford the free N10-Nvoc-Protected PBD-tripeptides.

Example 5(b): Photolytic Cleavage and MTT Assay Method

The same assay method as Example 3(b) was used. Cells at a
30 density of 5×10^4 cells/mL were continuously incubated with each member of the 27-mer library at a final concentration of 0.3 µM (6.6 µ moles/tube/ml). Aliquots of each of the compounds of the 27-member library were either left without UVA (365nm) exposure or were exposed to UVA (365nm) for 2h prior to their
35 addition to the cell suspension. Following addition of the compounds, the cells were transferred to 96-well microtitre plates, 10^4 cells per well, 8 wells per sample. Plates were incubated at 37°C in a humidified atmosphere containing 5% CO₂.

Following incubation of the plates for 4 days (to allow control cells to increase in number by 10-fold), 20 μ L of a 5mg/mL solution of MTT in phosphate-buffered saline was added to each well and the plates further incubated for 5 hours. The plates were then centrifuged for 5 minutes at 300g and the bulk of the medium was removed from the cell pellet, leaving 10-20 μ L per well. DMSO (200 μ L) was added to each well, and the samples agitated to ensure complete mixing. The optical density was then read at a wavelength of 550nm on a Titertek Multiscan ELISA plate reader and the dose-response curve constructed.

Results of *In vitro* cytotoxicity evaluation of the 27 member PBD library synthesised on 'crowns'

Compound	Amino Acid Sequence	% Control at 0.3 μ M
1	AAA	104
2	AAB	106
3	AAC	93.8
4	ABA	86.2
5	ABB	89.6
6	ABC	91.7
7	ACA	87.8
8	ACB	100.6
9	ACC	107
10	BAA	112
11	BAB	88.2
12	BAC	99.3
13	BBA	93.9
14	BBB	79.2
15	BBC	95
16	BCA	69.6
17	BCB	109
18	BCC	107.6
19	CAA	91.4
20	CAB	99.4
21	CAC	98.3
22	CBA	85.6
23	CBB	86.1
24	CBC	90.6
25	CCA	119
26	CCB	114
27	CCC	112
Benzyl DC-81	-	47.4

A=glycine

B=leucine

C=phenylalanine

Table 4: In vitro cytotoxicity of a 27 member PBD library synthesised on 'crowns'.

As before, these results demonstrate that varying the amino acid sequence affects the cytotoxicity of the PBDs.

Example 6: DNA Binding Assays

Labelling double-stranded oligonucleotides

Double-stranded oligonucleotides (10pmol/ μ L) were 5'-end labelled with [32 P]-ATP using T4 polynucleotide kinase and incubated for 30 minutes at 37°C. The labelled oligonucleotides were purified through a mini-prep Biorad™ spin column containing P6 Bio-gel™ (40-90 μ m).

On-bead screening assay

The beads were allowed to swell in DMF for approximately 1h prior to the binding experiment. Labelled double-stranded oligonucleotides were incubated with the beads to which compound was attached for 24h at 37°C. After 24h incubation the samples were resuspended in TE buffer (10mM tris, 1mM EDTA), spun and supernatant removed 3 to 4 times. On the final wash the pellet was resuspended in 1mL of EcoScint (Nat. Diagnostics, UK) scintillation fluid and counted on a Wallac 1400 scintillation counter.

Labelled oligonucleotides

Oligonucleotide 1(PuGPu) 5'- ACA CCT AIA **GAT** IAA ITC TI-3'
3'- TIT IIA TCT CTA CTT CAI AC-5'

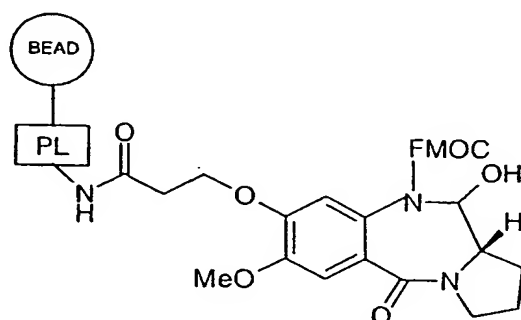
Oligonucleotide 2(PyGPy) 5'- ACA CCT AIT **GTT** IAA ITC TI-3'
3'- TIT IIA TCA CAA CTT CAI AC-5'

Oligonucleotide 3(PuIPu) 5'- ACA CCT AIA **IAT** IAA ITC TI-3'
3'- TIT IIA TCT CTA CTT CAI AC-5'

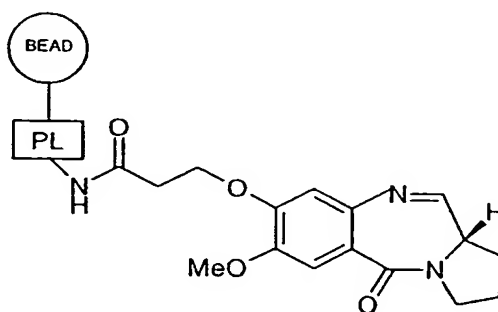
Oligonucleotide 1 contains the AGA sequence (highlighted in bold) which is the most preferred binding site for a PBD. Oligonucleotide 2 contains the TGT sequence which is the least

preferred binding site for a PBD. Oligonucleotide 3 contains the AIA sequence, to which a PBD should not bind owing to the lack of an NH₂ group on the inosine moiety.

5 Compound JGB-285 is N10-protected and should not bond
covalently to DNA:



Compound JGB-286 is a free C10-N11 imine moiety and is able to interact with DNA:



Compounds	Counts per minute (CPM)
JGB-285	0
JGB-286	50,394

Table 5: Binding of compounds JGB-285 and JGB-286 to Oligonucleotide 1 (counts corrected for background)

Double-stranded oligonucleotide	Counts per minute (CPM)
-	59.7
1	50,394
2	3,321
3	1,820

Table 6: Binding of JGB-286 to double-stranded

oligonucleotides 1, 2, and 3 (counts corrected for background)

These tests were also carried out using oligonucleotide 1 labelled with Rhodamine or Fluorescein instead of [³²P]-ATP (the labelled oligonucleotides are available from Genesis, Cambridge). The fluorescence was measured using a Tecan Spectrafluor Plus.

Compounds	Relative fluorescent units (RFU)
JGB-285	1,122
JGB-286	16,539

Table 7: Binding of compounds JGB-285 and JGB-286 to Rhodamine-labelled Oligonucleotide 1 (counts corrected for background).

Compounds	Relative fluorescent units (RFU)
JGB-285	1,217
JGB-286	42,355

Table 8: Binding of compounds JGB-285 and JGB-286 to Fluorescein-labelled Oligonucleotide 1 (counts corrected for background).

These results show that the PBD compound on-bead retains its ability to bind covalently to DNA - the protected PBD (JGB-285) did not bind at all, whereas the unprotected PBD (JGB-286) exhibited strong binding. More importantly, the unprotected PBD retained its selectivity for a PuGpu sequence, showing little binding activity towards the least preferred, and non-binding sites.

Example 7: Synthesis and Screening of Irori™ Glycine PBD Sublibrary (Figure 5)

Synthesis

Aminomethylated polystyrene resin 39 (5 g, 1.1 mmol/g loading) was suspended in DCE : CH₂Cl₂ (2:1, 102 ml) and dispensed equally into 289 Irori™ microkants. The resin was filtered and the kants were placed in a flask with DMF (70 ml) and shaken for 30 min.

A solution of Fmoc-glycine (4.91 g, 16.5 mmol), TBTU (5.3g, 16.5 mmol) and DIPEA (2.9 ml, 16.5 mmol) in DMF (100 ml) was added to the combined kans and shaking continued for 20 hours. Resin **40** was filtered and rinsed with CH₂Cl₂ (3 x 10 ml), MeOH (3 x 10 ml), Et₂O (3 x 10 ml) and dried *in vacuo*.

A solution of 20% Ac₂O, 30% pyridine in CH₂Cl₂ (200 ml) was added to the kans, which were shaken for 16 hours. The acetylated resin was filtered and washed with CH₂Cl₂ (3 x 10 ml), MeOH (3 x 10 ml), Et₂O (3 x 10 ml) and dried *in vacuo*.

A solution of 20% piperidine in DMF (200 ml) was added to the acetylated resin **40** and the reaction flask was shaken for 16 hours. Resin **41** was filtered and rinsed with CH₂Cl₂ (3 x 10 ml), MeOH (3 x 10 ml), Et₂O (3 x 10 ml) and dried *in vacuo*.

The kans were combined and sorted, using the Irori™ software into 17 flasks. Each flask contained a solution of an Fmoc-amino acid (0.97 mmol), TBTU (312 mg, 0.97 mmol) and DIPEA (170 ml, 0.97 mmol) in DMF (10 ml). [Fmoc-alanine (306 mg); Fmoc-asparagine (340 mg); Fmoc-aspartic (O^tBu) acid (391 mg); Fmoc-glutamine (357 mg); Fmoc-glutamic (O^tBu) acid (408 mg); Fmoc-glycine (289 mg); Fmoc-isoleucine (340 mg); Fmoc-leucine (340 mg); Fmoc(Boc)-lysine (357 mg); Fmoc-methionine (459 mg); Fmoc-phenylalanine (374 mg); Fmoc-proline (323 mg); Fmoc-serine (O^tBu) (374 mg); Fmoc-threonine (O^tBu) (391 mg); Fmoc(Boc)-tryptophan (510 mg); Fmoc-tyrosine (O^tBu) (442 mg); Fmoc-valine (323 mg)].

The flasks were shaken for 16 hours and each batch of 17 kans was filtered and rinsed with DMF (3 x 10 ml), CH₂Cl₂ (3 x 10 ml), MeOH (3 x 10 ml), Et₂O (3 x 10 ml) and dried *in vacuo* to give **42**.

A solution of 20% Ac₂O, 30% pyridine in CH₂Cl₂ (200 ml) was added to all 289 kans and the reaction flask was shaken for 16 hours. Acetylated resin was filtered and washed with CH₂Cl₂ (3 x 10 ml), MeOH (3 x 10 ml), Et₂O (3 x 10 ml) and dried *in vacuo*.

A solution of 20% piperidine in DMF (200 ml) was added to the acetylated resin and the vessel was shaken for 16 hours. Resin **43** was filtered and rinsed with CH_2Cl_2 (3 x 10 ml), MeOH (3 x 10 ml), Et_2O (3 x 10 ml) and dried *in vacuo*.

5

This process was repeated again to produce a library of trimers **45** and the final step was carried out simultaneously on all 289 kans.

10

The kans were placed in a flask with DMF (70 ml) and shaken for 30 min. A solution of Fmoc-PBD acid **7b** (Example 1/b) (9.2 g, 16.5 mmol), TBTU (5.3 g, 16.5 mmol) and DIPEA (2.9 ml, 16.5 mmol) in DMF (100 ml) was added to the kans and shaking continued for 20 hours. Resin **46** was filtered and rinsed with DMF (3 x 10 ml), CH_2Cl_2 (3 x 10 ml), MeOH (3 x 10 ml), Et_2O (3 x 10 ml) and dried *in vacuo*.

15

20

A solution of 2% triisopropylsilane in TFA (100 ml) was added to the kans suspended in CH_2Cl_2 (100 ml) and the kans were shaken for 16 hours. The kans were filtered and washed with CH_2Cl_2 (3 x 10 ml), MeOH (3 x 10 ml), Et_2O (3 x 10 ml) and dried *in vacuo*.

25

A solution of 20% piperidine in DMF (200 ml) was added to the kans and the reaction flask was shaken for 16 hours. The kans containing resin **47** were filtered and rinsed with CH_2Cl_2 (3 x 10 ml), MeOH (3 x 10 ml), Et_2O (3 x 10 ml), and dried *in vacuo*.

Screening

30

The resulting library was screened against a double stranded DNA sequence to determine which members of the library bound most strongly to the DNA test sequence.

The DNA sequence used was annealed fluorescein labelled:

35

Label-5'-ACACCTAAGATIAAITCTI-3'.

Approximately 10 mg of each library member was placed in a well of a 96 well plate and incubated with 5pmol/ μl annealed

fluorescein labelled double stranded DNA for 24 hours at 37°C. After 24 hours of incubation, each well was washed 4 times with TE buffer and the beads resuspended in 50 mL of TE or PBS.

- 5 The fluorescence of each well was measured using a Tecan Spectrafluor to determine which wells contained the most labelled DNA, and hence which compounds bound to the test DNA sequence most strongly.
- 10 Most active compounds from the library identified using the Irori™ software were found to be those with the following combinatorial chains: Gly-Gly-Gln-PBD; Gly-Pro-Iso-PBD; Gly-Thr-Asp-PBD; Gly-Leu-Val-PBD; Gly-Val-Asp-PBD; Gly-Val-Phen-PBD; Gly-Try-Asp-PBD; Gly-Lys-Ala-PBD; Gly-Gly-Asp-PBD; Gly-Gly-Pro-PBD.
- 15

Example 8: Synthesis of PBD - Glycine sublibrary (figures 6a, 6b, 6c)

Synthesis of Lysine-Glycine dimer 54 (Figure 6a)

- 20 Tentagel M NH₂ resin **48** (58 mg, 0.3 mmol/g loading) was weighed into 17 Alltech tubes (4 ml volume) and DMF (250 µl) was added to each tube, which were then shaken for 30 min.
- 25 A solution of Boc(Fmoc) lysine (416 mg, 0.88 mmol) in DMF (1.7 µl) and a solution of TBTU (285 mg, 0.88 mmol) and DIPEA (155 µl, 0.88 mmol) in DMF (3.4 ml) were equally dispensed into the tubes and shaking continued for 20 hours. Resin **49** was filtered and rinsed with DMF (3 x 2 ml), CH₂Cl₂ (3 x 2 ml), MeOH (3 x 2 ml) and dried *in vacuo*.
- 30 A solution of 20% Ac₂O, 30% pyridine in CH₂Cl₂ (500 µl) was added to each tube, and the tubes were shaken for 2 hours. The acetylated resin was filtered and washed with CH₂Cl₂ (3 x 2 ml), MeOH (3 x 2 ml), further CH₂Cl₂ (3 x 2 ml) and dried *in vacuo*.
- 35 A solution of 2% triisopropylsilane in TFA (250 µl) in CH₂Cl₂ (250 µl) was added to each tube, and the tubes were shaken for 2 hours. Resin **50** was filtered and washed with CH₂Cl₂ (3 x 2 ml), MeOH (3 x 2 ml), further CH₂Cl₂ (3 x 2 ml) and dried *in*

vacuo.

5 Resin 50 was suspended in CH_2Cl_2 (250 μl) and shaken for 30 min. An ice cold solution of allyl chloroformate (100 μl , 0.88 mmol) and 4-methylmorpholine (90 mg, 0.88 mmol) in CH_2Cl_2 (1.7 ml) was equally dispensed to each tube and shaken for 16 hours. Resin 51 was filtered and washed with CH_2Cl_2 (3 x 2 ml), MeOH (3 x 2 ml), further CH_2Cl_2 (3 x 2 ml) and dried *in vacuo*.

10 A solution of 20% piperidine in DMF (500 μl) was added to resin 51 and the tubes were shaken for 12 hours. Resin 52 was filtered and rinsed with DMF (3 x 2 ml), CH_2Cl_2 (3 x 2 ml), MeOH (3 x 2 ml) and dried *in vacuo*.

15 Resin 52 was suspended in DMF (250 ml) and shaken for 30 min.

20 A solution of Fmoc-glycine (264 mg, 0.88 mmol) in DMF (1.7 ml) and a solution of TBTU (285 mg, 0.88 mmol) and DIPEA (155 μl , 0.88 mmol) in DMF (3.4 ml) were equally dispensed to the tubes and shaking continued for 20 hours. Resin 53 was filtered and rinsed with DMF (3 x 2 ml), CH_2Cl_2 (3 x 2 ml), MeOH (3 x 2 ml) and dried *in vacuo*.

25 A solution of 20% Ac_2O , 30% pyridine in CH_2Cl_2 (500 μl) was added to resin 53 and the tubes were shaken for 2 hours. The acetylated resin was filtered and washed with CH_2Cl_2 (3 x 2 ml), MeOH (3 x 2 ml), further CH_2Cl_2 (3 x 2 ml) and dried *in vacuo*.

30 A solution of 20% piperidine in DMF (500 μl) was added to the acetylated resin and the tubes were shaken for 12 hours. Resin 54 was filtered and rinsed with DMF (3 x 2 ml), CH_2Cl_2 (3 x 2 ml), MeOH (3 x 2 ml) and dried *in vacuo*.

Synthesis of Glycine Sublibrary 61 (Figure 6b)

35 Coupling conditions

A solution of an Fmoc-amino acid (0.052 mmol) in DMF (100 μl) was added to each tube containing resin 54. [Fmoc-alanine (16 mg); Fmoc-asparagine (18 mg,); Fmoc-aspartic (O^tBu) acid (21

mg); Fmoc-glutamine (19 mg); Fmoc-glutamic (O^tBu) acid (22 mg); Fmoc-glycine (16 mg); Fmoc-isoleucine (18 mg); Fmoc-leucine (18 mg); Fmoc(Boc)-lysine (24 mg); Fmoc-methionine (19 mg); Fmoc-phenylalanine (20 mg); Fmoc-proline (18 mg); Fmoc-serine (^tBu) (20 mg); Fmoc-threonine (^tBu) (21 mg); Fmoc(Boc)-tryptophan (27 mg); Fmoc-tyrosine (^tBu) (24 mg); Fmoc-valine (18 mg)].

A solution of TBTU (285 mg, 0.88 mmol) and DIPEA (155 μ l, 0.88 mmol) in DMF (3.4 ml) was equally dispensed into the 17 tubes and shaking continued for 20 hours. Resin 55 was filtered and rinsed with DMF (3 x 2 ml), CH₂Cl₂ (3 x 2 ml), MeOH (3 x 2 ml) and dried *in vacuo*.

Acetylation conditions

A solution of 20% Ac₂O, 30% pyridine in CH₂Cl₂ (500 μ l) was added to resin 55 and the tubes were shaken for 2 hours. The acetylated resin was filtered and washed with CH₂Cl₂ (3 x 2 ml), MeOH (3 x 2 ml), further CH₂Cl₂ (3 x 2 ml) and dried *in vacuo*.

Deprotection conditions

A solution of 20% piperidine in DMF (500 μ l) was added to acetylated resin and the tubes were shaken for 2 hours. Resin 56 was filtered and rinsed with DMF (3 x 2 ml), CH₂Cl₂ (3 x 2 ml), MeOH (3 x 2 ml) and dried *in vacuo*.

Pool and Split method

The combined resin 56 was suspended in DCE : CH₂Cl₂ (2:1, 68 ml) in a round bottom flask, fitted with a sinter and was aspirated with nitrogen gas to ensure thorough mixing. The resin was re-dispensed into the 17 Alltech tubes, filtered and DMF (250 μ l) was added to each tube, followed by shaking for 30 minutes.

This cycle of coupling, acetylation, deprotection and pooling/splitting was repeated a further three times, until a library of 6-mer peptides 61 had been synthesised.

Synthesis of PBD-Glycine Sublibrary 64 (Figure 6c)

The resin 61 was suspended in CH₂Cl₂ (250 μ l) and the tubes were

shaken for 30 min. A solution of phenylsilane (876 μ l, 7.1 mmol) in CH_2Cl_2 (1.7 ml) was equally dispensed into each tube and the tubes were shaken for 10 min. A solution of tetrakis(triphenylphosphine)palladium (34 mg, 0.03 mmol) in CH_2Cl_2 (1.7 ml) was equally dispensed into each tube and the tubes were shaken for a further 10 min. Resin **65** was filtered and rinsed with CH_2Cl_2 (3 x 2 ml), MeOH (3 x 2 ml), further CH_2Cl_2 (3 x 2 ml) and dried *in vacuo*. This procedure was repeated once.

A solution of Fmoc PBD acid **7b** (Example 1b) (495 mg, 0.88 mmol) in DMF (3.4 ml) and a solution of TBTU (285 mg, 0.88 mmol) and DIPEA (155 μ l, 0.88 mmol) in DMF (3.4 ml) were equally dispensed to the suspension of 6-mer peptide resin **62** in DMF (250 μ l) and the tubes were shaken for 20 hours. Resin **63** was filtered and rinsed with DMF (3 x 2 ml), CH_2Cl_2 (3 x 2 ml), MeOH (3 x 2 ml) and dried *in vacuo*.

A solution of 2% triisopropylsilane in TFA (250 μ l) in CH_2Cl_2 (250 μ l) was added to resin **63** and the tubes were shaken for 2 hours. Resin was filtered and washed with CH_2Cl_2 (3 x 2 ml), MeOH (3 x 2 ml), further CH_2Cl_2 (3 x 2 ml) and dried *in vacuo*.

A solution of 20% piperidine in DMF (500 μ l) was added to the resin and the tubes were shaken for 2 hours. Resin **64** was filtered and rinsed with DMF (3 x 2 ml), CH_2Cl_2 (3 x 2 ml), MeOH (3 x 2 ml) and dried *in vacuo*.

The resulting sublibrary was screened against rhodamine labelled annealed double strand DNA with the sequence:
Label-5'-ACACCTAAGATIAAITCTI-3'

The sublibrary was mixed with the DNA sequence (5 pmol/mL) and incubated at 37°C for 24 hours with occasional mixing. After 24 hours, the sublibrary was washed 4 times with TE buffer pH 7.6 or PBS. To identify the beads to which most labelled DNA had bound, agarose gel slides were prepared as follows. ~ 500 mL of 0.25% sea plaque agarose was layered onto a clean

transparent slide and allowed to cool and set. The incubated beads were then mixed with another ~ 500 mL of 0.25% sea plaque agarose solution and layered onto the precoated slides, and allow to cool and set.

5

The reddest beads were identified by eye under a dissecting light microscope, and then retrieved by adding ~ 1 mL of water to dried agarose slide to enable their removal using a p10 gilson pipette with a fine tip. The removed beads were then placed into a 1 mL Eppendorf PCR tube ready for identification.

10

Identification

15

The identification of the sequences of the most active compounds was carried out using automated Edman degradation and reversed-phase HPLC.

20

Pulsed liquid-phase N-terminal sequencing was performed using an Applied Biosystems (ABI)477A automatic protein sequencer. The selected labelled beads were loaded onto a glass fibre disc which had previously been pre-cycled once. The disc was placed in the sequencer and pre-cycled once, then six cycles of Edman degradation were performed (Edman, P and Begg, G (1967) Eur. J. Biochem. 1, 80). The released phenylthiohydantoin (PTH-) amino acid derivatives were identified by reversed-phase HPLC analysis.

25

The eight most active compounds were those with the following sequences:

30

PBD-KGNNNN; PBD-KGTESF; PBD-KGMPMA; PBD-KGGGMM; PBD-KGKGAS;
PBD-KGANIA; PBD-KGMMGG; PBD-KGWYSP

35

Example 9: Synthesis of Split and Mix PBD Peptide Library 80 (Figure 7)

Aminomethylated polystyrene resin VHL 65 (3 g, 1.3 mmol/g loading) was suspended in DCE : CH₂Cl₂ (2:1, 102 ml) and

dispensed equally into 17 Alltech tubes (8 ml volume). The resin was filtered and DMF (1.5 ml) was added to each tube and shaken for 30 min.

5 Coupling conditions: (→66) - same as Example 8, but using a solution of an Fmoc-amino acid (0.69 mmol) in DMF (250 μ l) in each tube. (Fmoc-amino acids as example 8). The amounts of TBTU and DIPEA were increased in proportion to the amount of Fmoc-amino acid and were in 34ml of DMF.

10

Acetylation conditions: same as Example 8, but using 1.5 ml of CH_2Cl_2 containing the Ac_2O pyridine.

15

Deprotection conditions: (-67) same as Example 8, but using 1.5 ml solution of 20% piperidine in DMF.

Pool and Split method: - same as Example 8, but suspending resin in 102 ml of $\text{DCE}/\text{CH}_2\text{Cl}_2$.

20

The cycle was repeated a further four times, until a library of 5-mer peptides had been synthesised.

25

A solution of Boc(Fmoc) lysine (5.5 g, 11.7 mmol) in DMF (34 ml) and a solution of TBTU (3.76 g, 11.7 mmol) and DIPEA (2.04 ml, 11.7 mmol) in DMF (34 ml) were added to the suspension of 5-mer peptide resin 75 in DMF (17 ml) and the vessel was shaken for 20 hours. Resin 76 was filtered and rinsed with DMF (3 x 5 ml), CH_2Cl_2 (3 x 5 ml), MeOH (3 x 5 ml) and dried *in vacuo*.

30

A solution of 20% Ac_2O , 30% pyridine in CH_2Cl_2 (17 ml) was added to the resin and the vessel was shaken for 2 hours. The acetylated resin was filtered and washed with CH_2Cl_2 (3 x 5 ml), MeOH (3 x 5 ml), further CH_2Cl_2 (3 x 2 ml) and dried *in vacuo*.

35

A solution of 20% piperidine in DMF (17 ml) was added to the acetylated resin 76 and the vessel was shaken for 12 hours. Resin 77 was filtered and rinsed with DMF (3 x 5 ml), CH_2Cl_2 (3 x 5 ml), MeOH (3 x 5 ml) and dried *in vacuo*.

5 A solution of Fmoc PBD acid **7b** (Example 1b) (6.55 g, 11.7 mmol) in DMF (34 ml) and a solution of TBTU (3.76 g, 11.7 mmol) and DIPEA (2.04 ml, 11.7 mmol) in DMF (34 ml) were added to the suspension of 6-mer peptide resin **77** in DMF (17 ml) and the vessel was shaken for 20 hours. Resin **78** was filtered and rinsed with DMF (3 x 5 ml), CH₂Cl₂ (3 x 5 ml), MeOH (3 x 5 ml) and dried *in vacuo*.

10 A solution of 2% triisopropylsilane in TFA (17 ml) in CH₂Cl₂ (17 ml) was added to the resin and the vessel was shaken for 2 hours. Resin **79** was filtered and washed with CH₂Cl₂ (3 x 5 ml), MeOH (3 x 5 ml), further CH₂Cl₂ (3 x 2 ml) and dried *in vacuo*.

15 A solution of 20% piperidine in DMF (17 ml) was added to the resin **79** and the vessel was shaken for 2 hours. Resin **80** was filtered and rinsed with DMF (3 x 5 ml), CH₂Cl₂ (3 x 5 ml), MeOH (3 x 5 ml) and dried *in vacuo*.

20 Example 10: Synthesis of Glycine Sublibrary 87 (Figure 8)

Fmoc-aminoethyl photolinker NovaSyn TG resin **20** (30 mg, 0.23 mmol/g loading) was weighed into 17 Alltech tubes (4 ml volume) and a solution of 20% piperidine in DMF (250 ml) was added to each tube, which were shaken for 16 hours. Resin **21** was
25 filtered and rinsed with DMF (3 x 2 ml), CH₂Cl₂ (3 x 2 ml), MeOH (3 x 2 ml) and dried *in vacuo*. DMF (250 µl) was added to each tube and the tubes were shaken for 30 min.

30 Coupling Conditions:- Same as Example 8, but using solution of an Fmoc-amino acid (0.021 mmol) in DMF (150 µl) in each tube. The amounts of TBTU and DIPEA were increased in proportion to the amount of Fmoc-amino acid, and were in 1.7 ml DMF.

35 Acetylation Conditions: - Same as Example 8

Deprotection Conditions: - Same as Example 8

Pool and Split Method:- Same as Example 8

5 This cycle of coupling, acetylation, deprotection and pooling/splitting was repeated twice until a library of trimer peptides **83** had been synthesised, but after second repetition the resin was not pooled but kept as 17 separate sublibraries, allowing for the final amino acid to be known.

Coupling to Fmoc-glycine

10 Resin **83** was suspended in DMF (250 μ l) and shaken for 30 min. A solution of Fmoc-glycine (105 mg, 0.35 mmol) in DMF (1.7 ml) and a solution of TBTU (112 mg, 0.35 mmol) and DIPEA (68 μ l, 0.35 mmol) in DMF (1.7 ml) were dispensed equally to the tubes and shaking continued for 20 hours. Resin was filtered and
15 rinsed with DMF (3 x 2 ml), CH_2Cl_2 (3 x 2 ml), MeOH (3 x 2 ml) and dried *in vacuo*.

A solution of 20% Ac_2O , 30% pyridine in CH_2Cl_2 (500 μ l) was added to the tubes and were shaken for 2 hours. The acetylated
20 resin was filtered and washed with CH_2Cl_2 (3 x 2 ml), MeOH (3 x 2 ml), further CH_2Cl_2 (3 x 2 ml) and dried *in vacuo*.

A solution of 20% piperidine in DMF (500 μ l) was added to acetylated resin and the tubes were shaken for 2 hours. Resin
25 **84** was filtered and rinsed with DMF (3 x 2 ml), CH_2Cl_2 (3 x 2 ml), MeOH (3 x 2 ml) and dried *in vacuo*.

Coupling to Boc(Fmoc)-lysine

30 A solution of Boc(Fmoc)-lysine (165 mg, 0.35 mmol) in DMF (1.7 ml) and a solution of TBTU (112 mg, 0.35 mmol) and DIPEA (68 μ l, 0.35 mmol) in DMF (1.7 ml) were equally dispensed to the tubes and shaking continued for 20 hours. Resin was filtered and rinsed with DMF (3 x 2 ml), CH_2Cl_2 (3 x 2 ml), MeOH (3 x 2 ml) and dried *in vacuo*.

35

A solution of 20% Ac_2O , 30% pyridine in CH_2Cl_2 (500 μ l) was added to the tubes and were shaken for 2 hours. The acetylated resin was filtered and washed with CH_2Cl_2 (3 x 2 ml), MeOH (3 x

2 ml), additional CH_2Cl_2 (3 x 2 ml) and dried *in vacuo*.

5 A solution of 20% piperidine in DMF (500 μl) was added to acetylated resin and the tubes were shaken for 2 hours. Resin 85 was filtered and rinsed with DMF (3 x 2 ml), CH_2Cl_2 (3 x 2 ml), MeOH (3 x 2 ml) and dried *in vacuo*.

Coupling to PBD Capping Unit

10 Resin 85 was suspended in DMF (250 μl) and shaken for 30 min. A solution of Fmoc-PBD acid 7b (196 mg, 0.35 mmol) in DMF (1.7 ml) and a solution of TBTU (112 mg, 0.35 mmol) and DIPEA (68 μl , 0.35 mmol) in DMF (1.7 ml) were dispensed equally to the tubes and shaking continued for 20 hours. Resin 86 was
15 filtered and rinsed with DMF (3 x 2 ml), CH_2Cl_2 (3 x 2 ml), MeOH (3 x 2 ml) and dried *in vacuo*.

20 A solution of 2% triisopropylsilane in TFA (250 μl) and CH_2Cl_2 (250 ml) was added to the tubes which were shaken for 2 hours. Resin was filtered and rinsed with CH_2Cl_2 (3 x 2 ml), MeOH (3 x 2 ml) further CH_2Cl_2 (3 x 2 ml) and dried *in vacuo*.

25 A solution of 20% piperidine in DMF (500 μl) was added to the tubes which were shaken for 2 hours. Resin 87 was filtered and rinsed with DMF (3 x 2 ml), CH_2Cl_2 (3 x 2 ml), MeOH (3 x 2 ml) and dried *in vacuo*.

Example 11: Synthesis of Glycine Sublibrary 103 (Figure 9)

30 Aminomethylated resin 88 (30 mg, 0.97 mmol/g loading) was weighed into 17 Alltech tubes (4 ml volume). DMF (250 μl) was added to each tube and the tubes were shaken for 30 min.

35 Coupling Protocol:- Same as Example 8, but using a solution of an Fmoc-amino acid (0.087 mmol) in DMF (250 μl) in each tube. The amounts of TBTU and DIPEA were increased in proportion to the amount of Fmoc-amino acid.

Acetylation Condition:- Same as Example 8

Synthesis of Glycine Sublibrary 117 (Figure 10b)

Pool and Split Method: - Same as Example 8, starting with resin 109.

5

Coupling Conditions: - Same as Example 8 but using a solution of an Fmoc-amino acid (0.087 mmol) in DMF (250 μ l) in each tube.

10

Instead of TBTU and DIPEA a solution of diisopropylcarbodiimide (232 μ l, 1.48 mmol) and HOBt (200 mg, 1.48 mmol) in DMF (3.4 ml) was dispensed equally into the 17 tubes and shaken for 20 hours.

15

Acetylation Protocol: - Same as Example 8

Deprotection Protocol: - Same as Example 8

20

This cycle of pooling/splitting, coupling, acetylation and deprotection was repeated a further three times until a library of 6-mer peptides 117 had been synthesised, but after the third repetition the resin was not pooled but kept as 17 separate sublibraries, allowing for the final amino acid to be known.

25

Synthesis of Bis PBD-Glycine Sublibrary 123 (Figure 10C)

30

A solution of Boc(Fmoc)-lysine (695 mg, 1.48 mmol) in DMF (3.4 ml) and a solution of TBTU (480 mg, 1.48 mmol) and DIPEA (260 μ l, 1.48 mmol) in DMF (3.4 ml) were dispensed equally to resin 117 and shaking was continued for 20 hours. Resin 118 was filtered and rinsed with DMF (3 x 2 ml), CH₂Cl₂ (3 x 2 ml), MeOH (3 x 2 ml) and dried *in vacuo*.

35

A solution of 20% Ac₂O, 30% pyridine in CH₂Cl₂ (500 μ l) was added to resin 118 and the tubes were shaken for 2 hours. The acetylated resin was filtered and washed with CH₂Cl₂ (3 x 2 ml), MeOH (3 x 2 ml), further CH₂Cl₂ (3 x 2 ml) and dried *in vacuo*.

Resin 118 was suspended in CH₂Cl₂ (250 μ l) and the tubes were

shaken for 30 min. A solution of phenylsilane (1.5 ml, 11.9 mmol) in CH_2Cl_2 (3.4 ml) was dispensed equally into each tube and shaken for 10 min.

5 A solution of tetrakis(triphenylphosphine)palladium (57 mg, 0.05 mmol) in CH_2Cl_2 (3.4 ml) was dispensed equally into each tube and shaken for a further 10 min. Resin 119 was filtered and rinsed with CH_2Cl_2 (3 x 2 ml), MeOH (3 x 2 ml) and dried *in vacuo*. This procedure was repeated once.

10 A solution of 20% piperidine in DMF (500 μl) was added to resin 119 and the tubes were shaken for 2 hours. Resin 120 was filtered and rinsed with DMF (3 x 2 ml), CH_2Cl_2 (3 x 2 ml), MeOH (3 x 2 ml) and dried *in vacuo*.

15 Resin 120 was suspended in DMF (250 μl) and shaken for 30 min. A solution of Fmoc-PBD acid 7b (1.66 g, 2 x 1.48 mmol) in DMF (3.4 ml) and a solution of TBTU (960 mg, 2 x 1.48 mmol) and DIPEA (520 μl , 2 x 1.48 mmol) in DMF (3.4 ml) were equally
20 dispensed to the tubes and shaking continued for 20 hours. Resin 121 was filtered and rinsed with DMF (3 x 2 ml), CH_2Cl_2 (3 x 2 ml), MeOH (3 x 2 ml) and dried *in vacuo*.

25 A solution of 2% triisopropylsilane in TFA (250 μl) and CH_2Cl_2 (250 μl) was added to resin 122 and the tubes were shaken for 2 hours. Resin 122 was filtered and rinsed with CH_2Cl_2 (3 x 2 ml), MeOH (3 x 2 ml) further CH_2Cl_2 (3 x 2 ml) and dried *in vacuo*.

30 A solution of 20% piperidine in DMF (500 ml) was added to resin 122 and the tubes were shaken for 2 hours. Resin 123 was filtered and rinsed with DMF (3 x 2 ml), CH_2Cl_2 (3 x 2 ml), MeOH (3 x 2 ml) and dried *in vacuo*.

Example 13: Synthesis of Fmoc-Glu-OAll-Amino-Resin (171)
(Figure 11)

The TentaGel amine resin **169** (3.0 g, 0.8 mmol) in the form of
5 beads was allowed to swell for 2 hours in dry DMF (12 mL) and
shaken gently in a siliconised randomisation round bottom flask,
equipped with a sintered glass filter tube. The suspension was
filtered by suction from below. The beads were washed twice with
dry DMF as follows: DMF (20 mL) was added from the top of the
10 vessel (to wash down any resin adhering to the sides of the
vessel), nitrogen gas was gently bubbled from below through the
sintered glass for 2 minutes, and excess DMF was removed by
suction. A threefold molar excess of HOBt (7.7 mL, 0.3 M in DMF,
2.32 mmol) ("coupling reagent" was added to a threefold molar
15 excess of Fmoc-Glu-OAll (0.95 g, 2.32 mmol) and the resulting
solution was added to the resin. A threefold molar excess of
PyBOP (1.2 g, 2.32 mmol) and a threefold molar excess of DIPEA
(0.4 mL, 0.3 g, 2.32 mmol) in a minimal volume of DMF (1.2 mL)
were added to the reaction flask to initiate the coupling
20 reaction. The reaction flask was capped tightly and allowed to
shake gently for 1 hour at room temperature and excess coupling
reagent was removed by suction. The coupling procedure was
repeated once, from from the addition of coupling reagent to
ensure complete reaction. The resulting resin **171** was washed 6
25 times with DMF (4 x 20 mL), DCM (1 x 20 mL) and MeOH (1 x 20 mL),
on each occasion the resin was shaken for 2 minutes and then
filtered. The washing cycle was then repeated and the resin was
dried *in vacuo*.

30 Synthesis of Pentapeptide Library (172)

In order to cap any remaining free amino groups the resin (in the
randomisation flask), was suspended in a mixture of
Ac₂O/pyridine/DMF (0.2:0.3:0.5, 10 mL) and allowed to shake for 2
35 hours. The supernatant was removed by suction, the beads were
washed with DCM (1 x 10 mL), MeOH (1 x 10 mL) and again with DCM
(1 x 10 mL). The washing cycle was then repeated and the resin was
dried *in vacuo*.

5 The Fmoc protecting groups were removed by treating the resin with 50% piperidine in DMF (10.7 mL) whilst being shaken. After 10 minutes, the supernatant was removed by suction and fresh 50% piperidine in DMF (10.7 mL) was added and shaking continued. After another 10 minutes, the beads were washed 6 times with DMF (4 x 20 mL), DCM (1 x 20 mL) and MeOH (1 x 20 mL), on each occasion the resin was shaken for 2 minutes and then filtered. The washing cycle was then repeated and the resin was dried *in vacuo*.

10 The beads were suspended in an isopycnic mixture of DCE/DMF (2:1, 32 mL) and nitrogen gas was gently bubbled from below. Equal aliquots (470 μ L) of the suspension were added in sequence to each of the 17 Alltech tubes, previously marked with a letter corresponding to a specific amino acid. This was repeated 4 times.
15 The beads remaining in the randomisation flask were resuspended in the isopycnic mixture (32 mL) and the distribution process was then repeated twice.

20 The Glu-OAll-resin M1 (45.6 mmol) in each Alltech tube was allowed to swell in dry DMF (710 μ L) accompanied by gently shaking for 2 hours. Excess DMF was removed by suction on a vacuum manifold.

25 The resin was washed with dry DMF (1.18 mL) which was added from the top in order to wash down any resin adhering to the side of the Alltech tubes. The tubes were allowed to shake for 2 minutes and excess DMF was removed by suction on a vacuum manifold.

30 A threefold molar excess of HOBt (460 μ L, 0.3 M in DMF, 0.137 mmol) was added to a threefold molar excess of each Fmoc-amino-acid (0.137 mmol) and the resulting mixture was shaken for 10 minutes and then added to appropriate Alltech tube.

filtered using the vacuum manifold.

5 The beads were washed twice with dry DMF as follows: DMF (1.18 mL) was added from the top followed by gentle shaking for 2 minutes and excess DMF was removed by filtration on a vacuum manifold.

10 A threefold molar excess of HOBt (18 mg, 0.137 mmol) in a minimal volume of DMF (0.3 molar, 700 μ L) and threefold molar excess of Alloc-PBD 173 synthesised in an analagous manner to example 2 (55 mg, 0.137 mmol) in DMF were added to the resin.

15 A threefold molar excess of PyBOP (71 mg, 0.137 mmol) and threefold molar excess of DIPEA (24 μ L, 18 mg, 0.137 mmol) in a minimal volume of DMF (150 μ L), were added to the reaction tube to initiate the coupling reaction.

20 The reaction tube was capped tightly and allowed to shake gently for 16 hours at room temperature. Excess reagents were removed by filtration on a vacuum manifold.

25 The beads were washed 4 times with DMF (4 x 2 mL), DCM (1 x 2 mL) and MeOH (1 x 2 mL), on each occasion the resin 174 was shaken for 2 minutes and then filtered on a vacuum manifold. The washing cycle was then repeated and the resin was dried in vacuo.

Removal of Side Chain, Fmoc and Alloc Protecting Groups
(174-175)

30 The Boc and tBu protecting groups were removed by treating the PBD-peptide resin 174 (45.6 μ mol) in each Alltech tubes with a solution of TFA/triisopropylsilane/DCM (48:2:50, 800 μ L). The reaction tubes were allowed to shake for 30 minutes and excess reagents were removed by filtration on a vacuum manifold. The
35 procedure was repeated once and the beads were washed 3 times with DCM (1 x 2 mL), MeOH (1 x 2 mL) and again with DCM (1 x 2 mL), on each occasion the resin was shaken for 2 minutes and then filtered on a vacuum manifold. The washing cycle was then repeated and the

resin was dried *in vacuo*.

5 The resin was washed with DCM (5 x 2 mL) on each occasion the resin was shaken for 30 seconds and then filtered on a vacuum manifold.

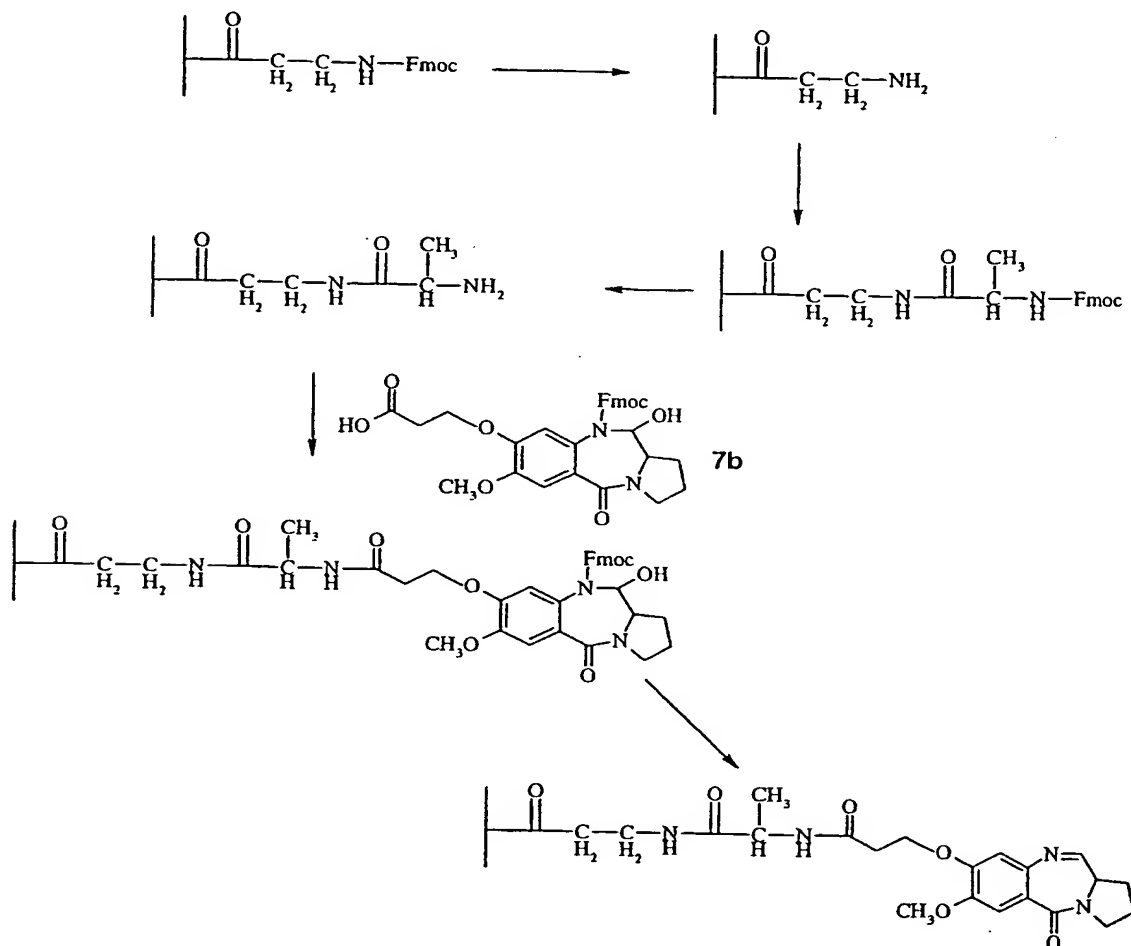
10 Alloc protecting groups were removed by treating the resin (45.6 μ mol) with a solution of phenylsilane (PhSiH_3 , 130 μ L, 0.118 g, 1.09 mmol) in DCM (300 μ L) and the resin was stirred manually. A solution of $\text{Pd}(\text{PPh}_3)_4$ (5.3 mg, 4.56 μ mol) in DCM (500 μ L) was added, and the Alltech tubes were shaken mechanically for 10 min. Excess reagents were removed by filtration on a vacuum manifold and the process was repeated once.

15 The peptide resin was washed with DCM (8 x 2 mL), on each occasion the resin was shaken for 30 seconds and then filtered on a vacuum manifold. The washing cycle was repeated and the resin was dried *in vacuo*.

20 The Fmoc groups were removed by treating the resin (45.6 μ mol) with 50% piperidine/DMF (800 μ L) during 2 hours on a shaker. The supernatant was removed by filtration on a vacuum manifold.

25 The beads 175 were washed 6 times with DMF (4 x 2 mL), DCM (1 x 2 mL) and MeOH (1 x 2 mL), on each occasion the resin was shaken for 2 minutes and then filtered on a vacuum manifold. The washing cycle was repeated twice and the resin was dried and stored *in vacuo*.

30

Example 14 - Cellulose Paper as a Laminar SupportA. Attachment of an Fmoc-PBD (7b) to Cellulose PaperModification of the cellulose paper - (General Method)

- 5 The required number of points were marked onto a square of cellulose paper, using a graphite pencil, before the dry paper was incubated with activated β -alanine solution (12 mL, [0.2M Fmoc- β -alanine activated with 0.24M DIC and 0.4M N-methylimidazole]), for 3hrs in a sealed vessel. The membrane was washed with DMF (3 x
- 10 50 mL for 3 minutes) and then treated with piperidine solution for 20 minutes (20% piperidine in DMF, 50. mL).

The membrane was washed with DMF (5 x 50 mL for 3 minutes) and MeOH (2 x 50 mL, for 3 minutes) and dried.

5 Fmoc- β -Alanine-OPfp solution (0.3M Fmoc- β Ala-OPfp in DMSO, 1 mL) was coupled to the pre-defined positions on the membrane. After 15 minutes, the coupling was repeated once again.

10 The membrane was then acetylated (2 minutes, face-down, in acetic anhydride solution A, 20mL [2% acetic anhydride in DMF], followed by 30 minutes, face-up in acetic anhydride solution B, 50 mL [2% acetic anhydride, 1% DIPEA in DMF], with shaking. After washing with DMF (3 x 50 mL for 3 minutes), the membrane was treated with piperidine solution for 20 minutes (20% piperidine in DMF, 50 mL), washed with DMF (5 x 50 mL for 3 minutes) and MeOH (2 x 50 mL for 3 minutes).

20 The membrane was stained with bromophenol blue solution (0.01% w/v bromophenol blue in methanol, 50 mL), washed with methanol for 3 minutes and dried.

Coupling of the Fmoc-PBD

25 A solution of Fmoc-PBD 7b, HOBT and DIC (0.3M, 1.5 mL) in NMP was spotted at the marked points on the membrane. This was left to couple for 1 hour and repeated (6x) until the blue colour of the spot was discharged. The membrane was washed once with DMF (50 mL) and incubated with acetic anhydride solution B, 50 mL for 30 minutes [2% acetic anhydride, 1% DIPEA in DMF].

30 The membrane was washed with DMF (5 x 50 mL) for 3 minutes, followed by methanol (2 x 50 mL) and dried.

N-Fmoc Deprotection

35 The Fmoc-protecting group was cleaved by treating the membrane with piperidine solution (20% in DMF, 20 mL) for 20 minutes.

The membrane was then washed with DMF (5 x 50 mL) for 3 minutes and MeOH (2 x 50 mL) for 3 minutes, stained with bromophenol blue solution (50 mL), washed with MeOH (50mL) for 3 minutes and dried.

[The bromophenol blue colouration was removed by destaining with piperidine solution, washing with DMF and MeOH and drying.]

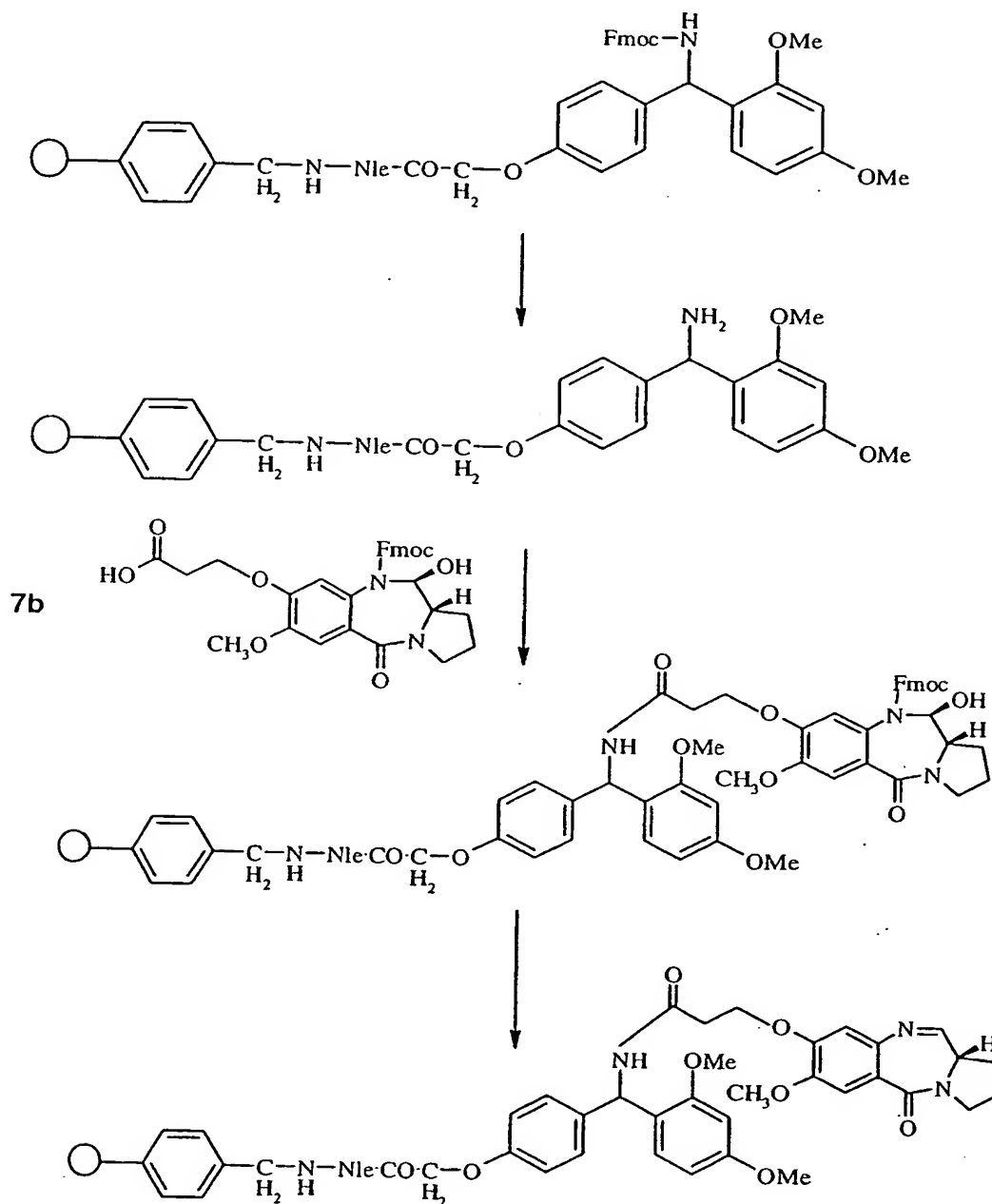
B. Attachment of Nvoc-PBD to Cellulose Paper

5 Coupling of the Nvoc-PBD (7a) (Example 1a)

The general method for modification and attachment of the Fmoc-PBD 7b was employed. The Nvoc-PBD (7a) was coupled as a 0.3 M solution for 30 minutes and repeated (4 x) until the bromophenol blue stain was discharged. The membrane was washed and dried as
10 previously described.

Deprotection of the Nvoc-protecting group

The membrane was incubated for several hours in a solution of DMSO containing 1% ethanolamine at a wavelength of 365 nm.
15 After washing with DMF (3 x 50 mL) for 3 minutes and MeOH (2 x 50 mL) for 3 minutes, the membrane was allowed to dry.

Example 15 - Synphase Crowns as Solid supportA. Attachment of Fmoc-PBD to Synphase crownsDeprotection of the Fmoc-protecting group

5

Two Fmoc-protected Rink-amide crowns (loading 7.7 mM/g), were placed in a scintillation vial and immersed in piperidine solution (20% in DMF, 2 mL) for 20 minutes, with shaking. After rinsing with DMF (3 x 2 mL) and DCM (3 x 2 mL), the

crowns were immersed in a solution of bromophenol blue (0.01% w/v bromophenol blue in MeOH, 5 mL), until equal colouration was achieved and the crowns were then rinsed in MeOH and dried.

5 Coupling of the Fmoc-PBD

A solution of Fmoc-PBD **7b** (17.18 mg, 3.08×10^{-5} mol), HOBT (8.32 mg, 6.16×10^{-5} mol) and diisopropylcarbodiimide (10.62 mL, 6.16×10^{-5} mol) in NMP (500 mL) was added to the scintillation vial with the crowns.

10

Coupling was monitored by the loss of the bromophenol blue colouration on the crowns. The reaction was allowed to proceed overnight and was repeated until complete colour loss had occurred. The crowns were then washed with DMF (3 x 2 mL), DCM (3 x 2 mL), MeOH (1 x 2 mL) and allowed to air-dry.

15

N-Fmoc Deprotection

A single crown was immersed in the piperidine solution (20% in DMF, 1 mL) for 20 minutes with shaking. After rinsing with DMF (3 x 1 mL), and DCM (3 x 1 mL), the crown was rinsed in MeOH (1 x 1 mL) and allowed to air-dry.

20

B. Attachment of Nvoc-PBD (7a) to Synphase crowns

Coupling of the Nvoc-PBD (7a)

25

The general method for modification and attachment of the Fmoc-PBD **7b** was employed. The Nvoc-PBD **7a** was coupled as a 0.62M solution overnight.

Deprotection of the Nvoc-protecting group

30

A single crown was immersed in DMSO containing 1% ethanolamine (2 mL) and irradiated at $\lambda=365$ nm for 2 hours, with shaking. After rinsing with DMF (3 x 1 mL), and DCM (3 x 1 mL), the crown was rinsed in MeOH (1 x 1 mL) and allowed to air-dry.

35

Example 16: Synthesis of a 175 membered Library utilising Synphase crowns as the solid support (Figure 12)

Preparation of the Crowns - N-Fmoc deprotection

175 Rink amide-handle O-series polystyrene crowns 124 were placed into two multipin blocks, arranged in a 8 x 12 format.

The crowns were immersed in the piperidine solution (20% in DMF, 50 mL) for 20 minutes, with shaking. After rinsing with DMF (3 x 50 mL) and DCM (3 x 50 mL), the crowns were immersed in a solution of bromophenol blue (0.01% w/v bromophenol blue in methanol, 50 mL), until equal staining was achieved and the deprotected crowns 125 were then rinsed in MeOH and dried.

Coupling the first amino acid residue

Amino Acid	M. Wt.	Reagent Mass (mg)	Solvent Vol. (mL)
Fmoc-Arg(Pbf)-OH	648.8	545.0	8.40
Fmoc-Gly-OH	297.3	254.8	8.40
Fmoc-Lys(Boc)-OH	468.5	393.5	8.40
Fmoc-Met-OH	371.5	312.1	8.40
Fmoc-Val-OH	339.4	285.1	8.40

Coupling Reagent (used for all couplings in example 16)

Reagent	M. Wt.	Reagent Mass	Solvent Vol
DIC	126.2	0.530g	42.0
HOBt	135.1	567.4 mg	42.0

Solutions of the amino acids and coupling reagents detailed above in NMP (200 mL) were dispensed into the deep-well microtiter plates. Couplings were monitored by the loss of the bromophenol blue staining on the crowns and were generally allowed to proceed overnight. The crowns 126 were then washed in DMF (3 x 50mL), DCM (3 x 50 mL), and allowed to air-dry.

N-Fmoc deprotection

The crowns 126 were immersed in piperidine solution (20% in DMF, 50 mL) for 20 minutes, with shaking. After rinsing with DMF (3 x 50 mL) and DCM (3 x 50 mL), the deprotected crowns 127 were immersed in a solution of bromophenol blue (0.01% w/v bromophenol

blue in methanol, 50 mL), until equal staining was achieved and then rinsed in MeOH and dried.

Coupling the second amino acid residue

Amino Acid	M.Wt.	Reagent Mass (mg)	Solvent Vol. (mL)
Fmoc-Arg(Pbf)-OH	648.8	389.3	6.0
Fmoc-Gly-OH	297.3	182.0	6.0
Fmoc-Lys(Boc)-OH	468.5	281.1	6.0
Fmoc-Gln-OH	368.4	221.0	6.0
Fmoc-His(Trt)-OH	619.7	379.4	6.0
Fmoc-Leu-OH	353.4	212.0	6.0
Fmoc-Tyr(2-ClTrt)-OH	680.2	408.1	6.0

Solutions of the amino acids and coupling reagents detailed above (in NMP, 200 mL) were dispensed into the deep-well microtiter plates. Couplings were monitored by the loss of the bromophenol blue staining on the crowns 127 and were generally allowed to proceed overnight. The coupled crowns 128 were then washed in DMF (3 x 50mL), DCM (3 x 50mL) and were allowed to air dry.

N-Fmoc deprotection was carried as above, to give crowns 129.

Coupling the third amino acid residue

Amino Acid	M.Wt.	Reagent Mass (mg)	Solvent Vol. (mL)
Fmoc-Arg(Pbf)-OH	648.8	545.0	8.40
Fmoc-Lys(Boc)-OH	468.5	393.5	8.40
Fmoc-Gly-OH	297.3	254.8	8.40
Fmoc-Gln-OH	368.4	309.5	8.40
Fmoc-Trp-OH	426.5	358.3	8.40

Solutions of the amino acids and coupling reagents detailed above (in NMP, 200 mL) were dispensed into the deep-well microtiter plates. Couplings were monitored by the loss of the bromophenol blue staining on the crowns and were generally allowed to proceed overnight. The coupled crowns 130 were then washed in DMF (3 x 50mL), DCM (3 x 50 mL) and allowed to air dry overnight.

N-Fmoc deprotection was carried out as above to give crowns 131.

Coupling the Fmoc-PBD

Reagent	M. Wt.	Reagent Mass (g)	Solvent Vol. (mL)
Fmoc-PBD 7b	558	2.346	42.0

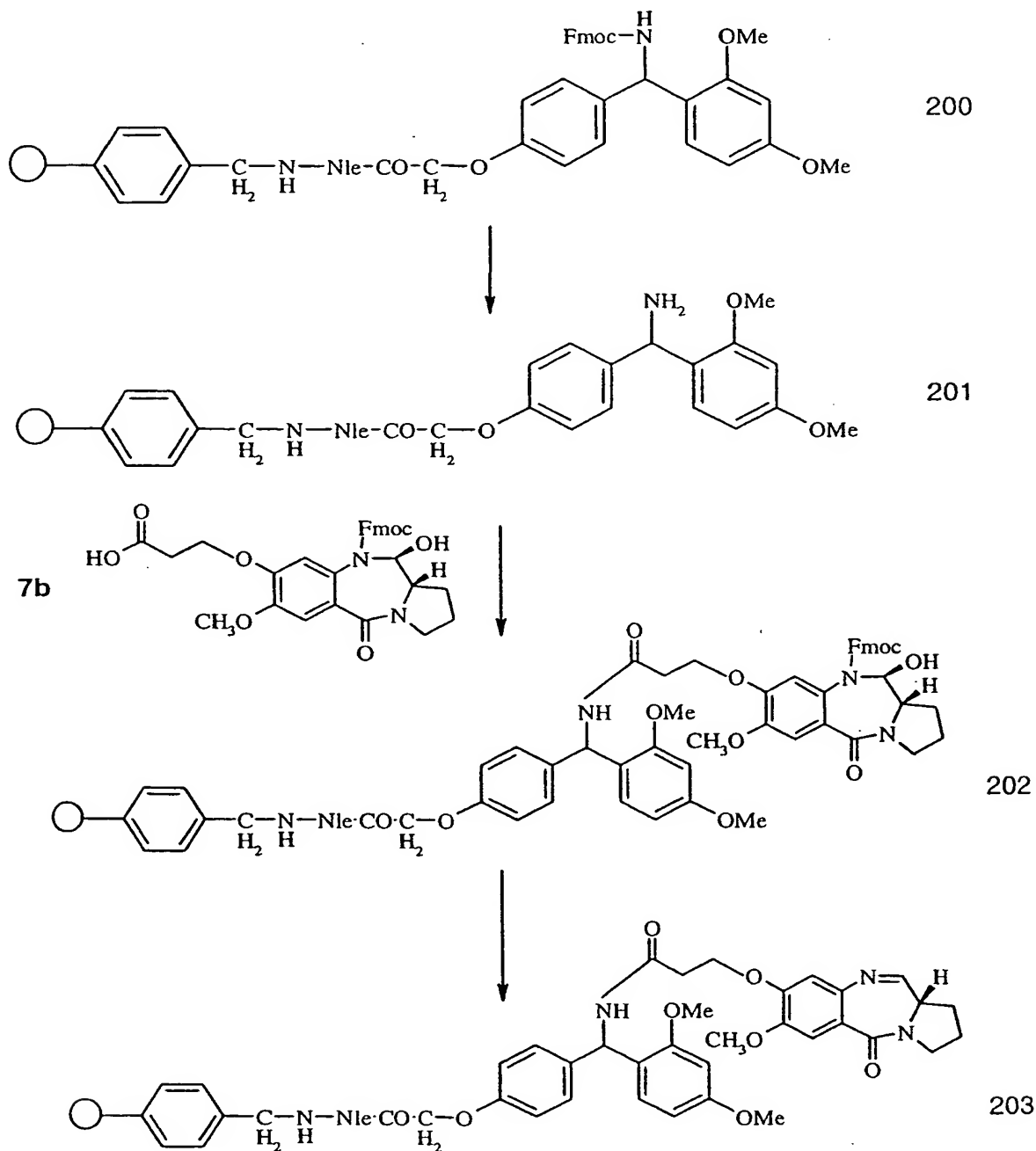
5 Fmoc-PBD **7b** (2.346g in 42 ml NHP) and the coupling reagents
detailed above (in NMP, 200 mL) were dispensed into the deep-well
microtiter plates. Couplings were monitored by the loss of the
bromophenol blue staining on the crowns **132** and repeated until
complete colour loss had occurred. The crowns were then washed
10 in DMF (3 x 50 mL), DCM (3 x 50 mL) and allowed to air dry.

N-Fmoc deprotection

The crowns **132** were immersed in piperidine solution (20% in DMF,
50 mL) for 20 minutes, with shaking. After rinsing with DMF (3
15 x 50 mL) and DCM (3 x 50 mL), the crowns **133** were allowed to air
dry.

Example 17: Use of Rink-amide Resin as Solid SupportA. Attachment of Fmoc-PBD (7b) to Rink-amide ResinPreparation of the resin - N-Fmoc deprotection

Fmoc-rink amide resin 200 (50 mg, loading 0.63 mmol/g) was



5 suspended in DCM:DMF (1:1, 1 mL) and shaken for 2 minutes. The

resulting solution was removed under vacuum and the resin resuspended in DMF (1 mL) and shaken for 5 minutes.

The DMF was drained from the resin and the Fmoc-group was removed by treatment with piperidine solution (20% in DMF, 1 mL) for 20 minutes. Excess piperidine solution was removed by suction and the deprotected resin **201** was washed with DMF and DCM and allowed to dry.

Coupling of the Fmoc-PBD

A solution of Fmoc-PBD propanoic acid **7b** (70.3 mg, 1.26×10^{-4} mol) diisopropylcarbodiimide (20.1 mL 1.26×10^{-4} mol) and HOBt (17.02 mg, 1.26×10^{-4} mol) in DMF (0.5 mL) was added to the resin **201**, and the resulting slurry shaken overnight.

The solution was drained and the coupled resin **202** washed with DMF and DCM. The resin was resuspended in DMF/DCM (DCM) (1 mL) and split into two portions. The solutions were removed from both tubes and the resin in the first tube was washed with DCM, methanol and dried under vacuum overnight.

N-Fmoc deprotection

The resin **202** in the second tube was suspended in piperidine solution (20% in DMF, 0.5 mL) and shaken for 20 minutes. Excess piperidine solution was drained and the deprotected resin **203** washed with DMF, DCM and methanol, and dried under vacuum overnight.

B. Attachment of Nvoc-PBD (7a)

Preparation of the resin - N-Fmoc deprotection

Coupling of the Nvoc-PBD (7a)

The general method for modification and attachment of the Fmoc-PBD **7b** was employed. The Nvoc-PBD **7a** was coupled as a 0.25M solution in DMF.

Example 18: Attachment of Fmoc-PBD (7b) to Aminoethyl Photolinker AM Resin (210)

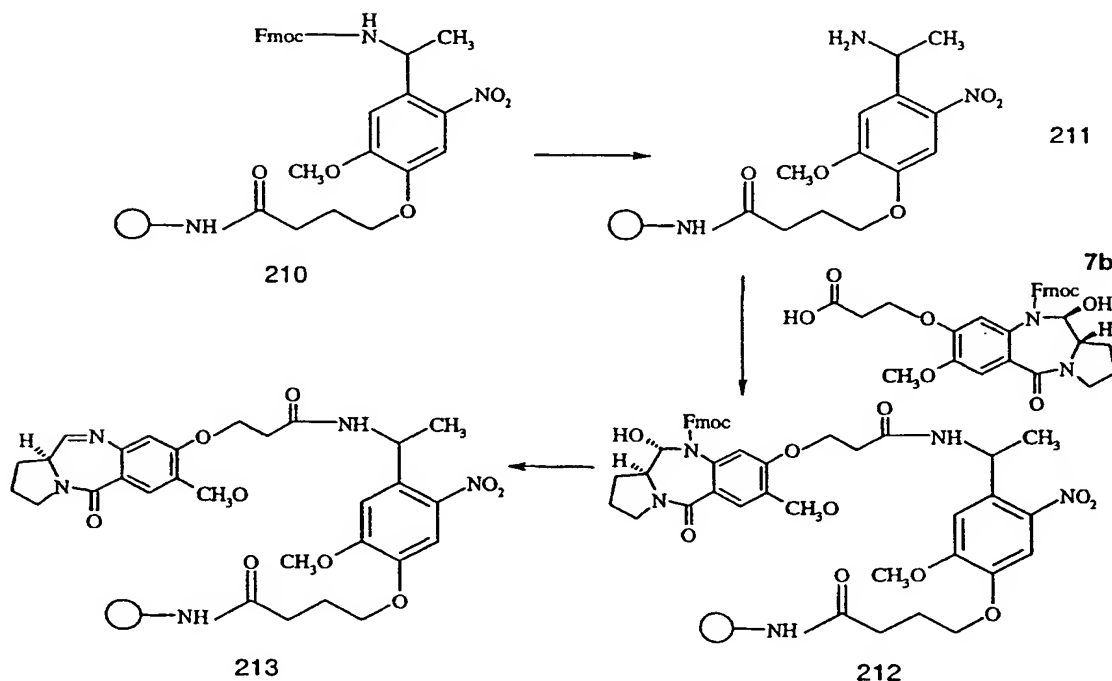
Preparation of the Resin - N-Fmoc deprotection

5 Aminoethyl photolinker AM resin **210** (50 mg, 0.21 mmol/g) was placed into two Alltech tubes. The resin was suspended in NMP (2 mL) and swelled for 2 hours.

10 After draining excess solvent, the deprotected resin **211** was suspended in piperidine solution (20% in DMF, 0.5 mL) and shaken for 20 minutes. Excess solvent was drained and the resin washed with DMF, DCM and NMP.

Coupling Protocol of Fmoc-PBD (7b)

15 A solution of Fmoc-PBD propanoic acid **7b** (32.22 mg, 5.78×10^{-5}



mol, 5.5 equivalents), PyBop (11.4 mg, 2.2×10^{-5} mol) and DIPEA (0.05 mL) in NMP (0.5 mL) was added to the resin **211** which was shaken for 1 hour.

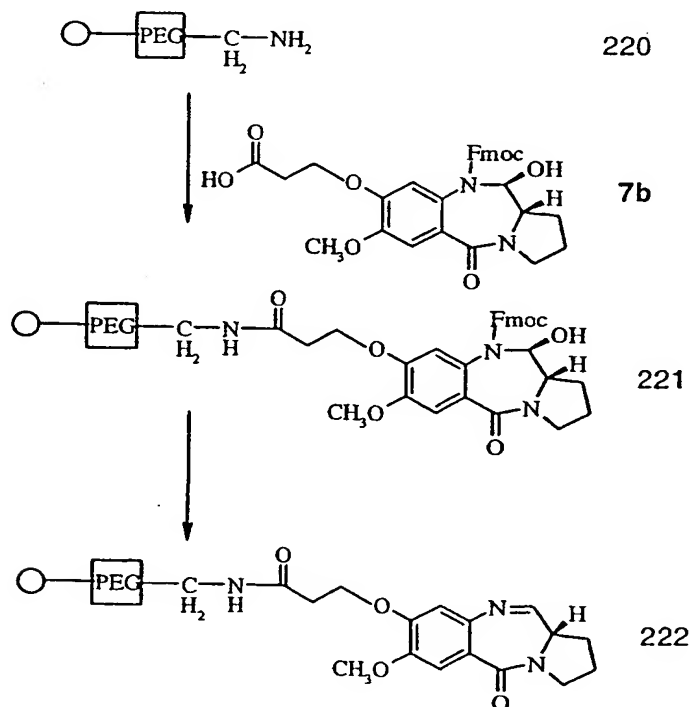
After draining, the coupled resin **212** was washed with DCM (5 x 1 mL), methanol (5 x 1 mL) and NMP (5 x 1 mL). The coupling reaction was repeated 3 times. The resin in the first tube was then washed with DCM (3 x 1 mL), methanol (3 x 1 mL) and water (3 x 1 mL) and dried *in vacuo* overnight.

N-Fmoc deprotection

The Fmoc-PBD resin **212** in the second Alltech tube was re-suspended in piperidine solution (20% in DMF, 1 mL) and shaken for 20 minutes. After draining, the deprotected resin **213** was washed with NMP (3 x 1 mL), DCM (3 x 1 mL), methanol (3 x 1 mL) and finally water (3 x 1 mL) and dried *in vacuo* overnight.

Example 19: Tentagel-Amino Resin

A. Attachment of Fmoc-PBD (7b) to Novasyn Tentagel Amino Resin (220)



Preparation of the resin

Novasyn Tentagel amino resin **220** (50 mg, loading 0.28 mmol/g), was placed into two Alltech tubes and suspended in DCM:DMF (1:1, 1 mL)

and shaken for 2 minutes. Excess solvent was removed by suction and the resin resuspended in DMF (1 mL), shaken for 5 minutes and then drained.

5 Coupling of Fmoc-PBD (7b)

A solution of Fmoc-PBD propanoic acid **7b** (23.44 mg, 4.2×10^{-5} mol, 3 x excess), TBTU (13.4 mg, 4.2×10^{-5} mol) and DIPEA (7.3 mL, 4.2×10^{-5} mol) in DMF (500 mL) was added to the resin **220** and the resulting slurry shaken overnight.

10

The solution was drained and the resin washed with DMF and DCM. The coupling was repeated a second time and the coupled resin **221** washed as before. The resin in the first tube was washed further with methanol and dried *in vacuo*, overnight.

15

Fmoc deprotection

The resin **221** in the second tube was suspended in piperidine solution (20% in DMF, 0.5 mL) and shaken for 20 minutes. This solution was drained and the deprotected resin **222** washed with DMF, DCM and methanol, and dried under vacuum overnight.

20

B. Attachment of Nvoc-PBD (7a) to Tentagel Resin

Coupling of the Nvoc-PBD (7a)

The general method for modification and attachment of the Fmoc-PBD **7b** was employed. The Nvoc-PBD **7a** was coupled as a 0.085M solution.

25

Deprotection of the Nvoc-protecting group

The resin was incubated for several hours in a solution of DMSO containing 1% ethanolamine at a wavelength of 365 nm. After washing with DMF (3 x 50 mL) for 3 minutes and MeOH (2 x 50 mL) for 3 minutes, the membrane was allowed to dry.

30

Example 20: Synthesis of an PBD-Oligocarbamate Sequence (Figure 13)

35

Preparation of the resin (N-Fmoc Deprotection)

Fmoc-Rink amide resin **134** (53.43 mg, loading 0.63 mmol/g, 3.37

x 10⁻⁵ mol) was suspended in DCM:DMF (1:1, 1 mL) and shaken for 2 minutes. Excess solvent was removed by suction, the resin resuspended in DMF (1 mL) and shaken for 5 minutes.

- 5 Excess DMF was removed by suction and the Fmoc-group was removed by treating with piperidine solution (20% in DMF, 1 mL) for 20 minutes. The resin 135 was then washed with DMF and DCM.

Coupling Protocol

	Fmoc aminoalkyl carbonate	M.Wt	No of moles 4x excess	Mass required/ mg
10	1 Fmoc-Lys(Boc) ^c	619	1.35 x 10 ⁻⁴	83.6
	2 Fmoc-Phe ^c	539	1.35 x 10 ⁻⁴	72.8
	3 Fmoc-Ser(^t Bu) ^c	535	1.35 x 10 ⁻⁴	72.2
	4 Fmoc-Gly ^c	448	1.35 x 10 ⁻⁴	60.5
15	5 Fmoc-Tyr(^t Bu) ^c	610	1.35 x 10 ⁻⁴	82.4
	6 Fmoc-Ile ^c	504	1.35 x 10 ⁻⁴	68.0
	7 Fmoc-Tyr(^t Bu) ^c	610	1.35 x 10 ⁻⁴	82.4

- 20 A solution of the required 4-nitrophenyl Fmoc-aminoalkyl carbonate, HOBt (36.3 mg) and DIPEA (11 ml), in NMP (200 ml) was added to the free amino resin 135 and allowed to couple for 4 hours. After draining, the resin 136 was washed with NMP and DCM and dried under vacuum.

- 25 Deprotection and coupling cycles were repeated until the aminoalkyl sequence was complete (137 - 148).

- 30 Fmoc-protected PBD (7b) (75.3mg), HOBt (36.3mg) and diisopropylcarbodiimide (21.5ml) were dissolved in NMP (300ml) and added to the free amino resin 148. The slurry was shaken for 24 hours, drained and washed with NMP, DCM and dried overnight, under vacuum to give PBD resin 149. This was deprotected as above to give unprotected PBD resin 150.

Cleavage from the resin

The resins (149 and 150) were treated with a solution of 95% TFA :2.5% water :2.5% triisopropylsilane (1 mL) at RT for 2 hours. The resins were removed by filtration, washed with a small amount of TFA and DCM. The solvent was removed in vacuo

and the residue dissolved in acetonitrile : water 1:2 and lyophilised twice.

Example 21: Synthesis of a 289 member library using 4-nitrophenyl N-Fmoc aminoalkyl carbonates (Figure 14)

Formation of N-(9-fluorenylmethoxycarbonyl) amino alcohols.
(General method of Cho et al : Synthesis and Screening of
Linear and Cyclic Oligocarbamate Libraries. Discovery of High
Affinity Ligands for GPIIb/IIIa, J. Am. Chem. Soc., (1998),
120, 7706-7718., C. Y. Cho, R. S. Youngquist, S. J. Paikoff,
M. H. Beresini, A. R. Herbert, L. T. Berleau, C. W. Liu, D. E.
Wemmer, T. Keough and P. G. Schultz.)

The appropriate N-Fmoc-protected amino acid (10 mmol) and dimethoxyethane were stirred under nitrogen, in an ice/salt bath. N-methylmorpholine (1.11 mL, 10 mmol) and isobutylchloroformate (1.36 mL, 10 mmol) was added to the solution.

After stirring for 1 min, under nitrogen, the solid was removed by filtration and sodium borohydride (570 mg, 15 mmol) in water (20 mL), was added to the filtrate. Additional water (150 mL) was added after 20 minutes, and the solution allowed to stir for 1 hour at room temp.

The precipitated product was filtered and washed with a small amount of water followed by hexane. The solid was redissolved in ethyl acetate, dried using MgSO₄ and the solvent removed *in vacuo*.

For those amino alcohol derivatives that did not precipitate from solution, the solution was extracted with ethyl acetate (5 x 300 mL). The organic extract was dried, over MgSO₄ and the solvent removed *in vacuo*.

Fmoc-amino alcohols were used without further purification - the data for the amino alcohols are shown in Appendix 1

Formation of 4-nitrophenyl N-(9-fluorenylmethoxycarbonyl) - aminoalkyl carbonates (General method of Cho et al.)

To the appropriate N-Fmoc-amino alcohol (10 mmol) was added pyridine (11 mmol) and DCM (50 mL). A solution of 4-nitrophenyl chloroformate (11 mmol) in DCM (10mL) was added dropwise to the reaction mixture. The mixture was stirred for at least 24 hours.

The mixture was diluted with DCM (100 mL) and washed with 1.0 M sodium bisulfate (3 x 75 mL) and 1.0 M sodium bicarbonate (10 x 100 mL). The organic layer was dried, using MgSO₄, and the solvent removed *in vacuo*.

The crude product was purified using silica gel chromatography (9:1 DCM:hexane DCM). The data for the amino alkyl nitrophenyl carbonate is shown in Appendix 2.

Library Synthesis

Preparation of the resin

Rink amide MBHA resin **151** (8.67 g, loading 0.54 mMol/g) was suspended in a solution of DCM/DMF (3:1) and distributed between 289 Irori MicroKan reactors containing RF tags.

The Kans were immersed in a solution of DCM/DMF (1:1, 300mL) and shaken vigorously for 1 min. The solvent was removed under vacuum and the Kans were re-immersed in DMF and shaken vigorously for 10 minutes.

The DMF was drained and the Fmoc-group was removed by adding piperidine solution (20% in DMF, 300 mL) to the Kans followed by shaking for 5 minutes and draining. Additional piperidine solution was added (20% in DMF) and the Kans were shaken for 1 hour. The Kans with deprotected resin **152** were drained and washed with DMF (6 x 300 mL) and DCM (3 x 300 mL).

Coupling of the first 4-nitrophenyl fluorenylmethoxy aminoalkyl carbonate

A solution of Fmoc-Valine^c (6.88g, 4.86×10^{-5} mol, 3 x excess), HOBT (3.795g, 9.72×10^{-5} mol) and DIPEA (1.22 mL, 2.43×10^{-5} mol) in NMP (300 mL) was added to the Kans and allowed to couple for 4 hours.

After draining the Kans with coupled resin **153** were washed with NMP (3 x 300 mL) and DCM (3 x 300 mL).

Fmoc-deprotection

The Fmoc-protecting group was removed by adding piperidine solution to the Kans (20% in DMF, 300mL) followed by shaking for 1 hour. The Kans with deprotected resin **154** were drained and washed with DMF (3 x 300 mL) and DCM (3 x 300 mL).

Coupling of the second 4-nitrophenyl fluorenylmethoxy aminoalkyl carbonate

Fmoc-Amino alkyl nitrophenyl carbonate	Mass (mg)
Ala	382
Asn	417
Asp(O ^t Bu)	467
Arg(Pbf)	673
Gln	428
Glu(O ^t Bu)	449
Gly	370
Ile	416
Leu	416
Lys(Boc)	511
Met	430
Phe	445
Pro	477
Ser(^t Bu)	442
Thr(^t Bu)	453
Trp	477
Tyr(^t Bu)	504

The Kans with the resin **154** were sorted using their Rf tags into 17 flasks containing solutions of the appropriate 4-nitrophenyl Fmoc- amino alkyl carbonate (8.262×10^{-4} mol), HOBT (223 mg, 1.6524×10^{-3} mol) and DIPEA (71.4 mL, 4.131×10^{-4} mol) in NMP (15 mL). The Kans were agitated four 4 hours, drained and washed with

NMP (3 x 100 mL) and DCM (3 x 100 mL), and then contained coupled resin 155).

N-Fmoc Deprotection

The 289 Kans with coupled resin 155 were pooled in one flask and the Fmoc-protecting group was removed by treatment with piperidine solution (20% in DMF, 300 mL) for 1 hour. The Kans with deprotected resin 156 were drained and washed with DMF (3 x 300 mL) and DCM (3 x 300 mL).

Coupling of the third 4-nitrophenyl fluorenylmethoxy aminoalkyl carbonate

Fmoc-Amino alkyl nitrophenyl carbonate	Mass (mg)
Ala	382
Asn	417
Asp(O ^t Bu)	467
Arg(Pbf)	673
Gln	428
Glu(O ^t Bu)	449
Gly	370
Ile	416
Leu	416
Lys(Boc)	511
Met	430
Phe	445
Pro	477
Ser(^t Bu)	442
Thr(^t Bu)	453
Trp	477
Tyr(^t Bu)	504

The Kans with resin 156 were sorted via their Rf tags into 17 flasks containing solutions of the appropriate 4-nitrophenyl Fmoc-amino alkyl carbonate (8.262×10^{-4} mol), HOBt (223mg, 1.6524×10^{-3} mol) and DIPEA (71.4ml, 4.131×10^{-4} mol) in NMP (15mL). The resin in the Kans was allowed to couple for 4 hours to form coupled resin 157. After draining, the Kans were washed with NMP (3 x 100mL) and DCM (3 x 100mL).

N-Fmoc Deprotection was carried out as above to give resin 158 in the Kans.

Coupling of the Fmoc-PBD (7b)

A solution of Fmoc-PBD (7b) (7.84 g, 1.40×10^{-2} mol, 3x excess), HOBt (1.896 g, 1.40×10^{-2} mol) and diisopropylcarbodiimide (2.18 mL, 1.40×10^{-2} mol) in NMP (300 mL) was added to the 289 Kans with resin 158. The Kans were stirred vigorously for 24 hours and drained. The Kans which then contained coupled resin 159 were washed with NMP (3 x 300mL) and DCM (3 x 300 mL).

N-Fmoc deprotection was carried out as before to give resin 160 in the Kans.

Example 22: Synthesis of a 27 member library using Peptoids (Irori method) (Figure 15)Preparation of the resin

Rink amide MBHA resin 151 (810mg, loading 0.54mMol/g) was suspended in a solution of DCM/DMF (3: 1, 5.4mL) and distributed between 27 Irori MicroKan reactors containing RF tags.

The Kans were immersed in a solution of DCM/DMF (1:1, 50mL) and shaken vigorously for 1 min. This solution was removed under vacuum and the Kans were re-immersed in DMF and shaken vigorously for 10 minutes.

The DMF was drained and the Fmoc-group of resin 151 was removed by adding piperidine solution (20% in DMF, 50mL) to the Kans. This was shaken for 5 minutes and drained. Further piperidine solution was added (20% in DMF, 50mL) and the Kans shaken for 1 hour. The Kans with deprotected resin 152 were drained and washed with DMF (6 x 50mL).

Acylation Step

Bromoacetic acid solution (0.6M in DMF, 60mL) was added to the Kans with resin 152, followed by diisopropylcarbodiimide solution (3.2M in DMF, 14.1mL). This was shaken for 2 hours at

room temperature, drained and repeated. The Kans with bromoacetamide resin **161** were then washed with DMF (2 x 50mL) and DMSO (1 x 50mL).

5 Displacement step

The 27 Kans were sorted via their RF tags into 3 flasks. The first set were suspended in aq. methylamine solution (40% w/v, 20mL), the second set in piperonylamine solution (2M in DMSO, 20mL) and the third set in 2-methoxyethylamine (2M in DMSO, 20mL). These were shaken vigorously for 4 hours and then drained. The Kans with amino-coupled resin **162** were washed with DMSO (2 x 20mL) and DMF (1x 20ml).

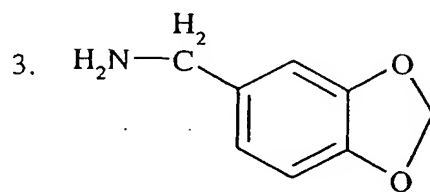
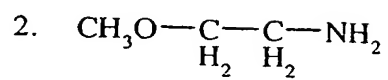
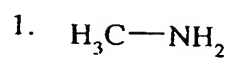
15 The acylation and displacement steps were repeated twice to give resin **166** (which is a library of 27 resins with all combinations of 3 values for R_1 , R_2 and R_3).

Coupling of the Fmoc-PBD (7b)

20 The 27 Kans with the peptoid bearing resin **166** were re-combined and a solution of the Fmoc-PBD (7b) (976.3mg, 1.75mmol, 4 x excess), HOBt (236.4mg, 1.75mmol) and diisopropylcarbodiimide (22.08 mg, 1.75mmol) in DMF (20 mL) added. This was stirred vigorously for 24 hours and drained. The Kans with the Fmoc-PBD coupled resin **167** were washed with DMF (3x 50mL) and DCM (3 x 50mL).

N-Fmoc Deprotection

30 The Fmoc-protecting group was removed from the PBD resin **167** by adding piperidine solution (20% in DMF, 50mL) to the Kans. This was shaken for 5 minutes and drained. Further piperidine solution was added (20% in DMF, 50mL) and the Kans shaken for 1 hour. The Kans with the deprotected resin **168** were drained and washed with DMF (3 x 50mL), DCM (3 x 50mL), methanol (2 x 50mL) and dried overnight under vacuum.

Amines used in library

APPENDIX 1

Data for Amino Alcohols incl yield (%) &NMRAla:-74%

5 ^1H NMR (270 MHz, d_6 -acetone): δ 7.88 (d, 2, J = 7.1 Hz), 7.71 (d, 2, J = 7.3 Hz), 7.31-7.42 (dt, 4), 7.09 (d, 1, J = 8.1 Hz), 4.28 (m, 4), 3.54 (m, 1), 3.36 (m, 1), 3.26 (m, 1), 1.04 (d, 3, J = 6.6 Hz).

^{13}C NMR (67.8 MHz, d_6 -acetone): δ 155.5, 143.9 140.7 127.5, 127.0, .1, 120.0, 65.1 64.4, 48.4 46.7 17.3

Arg(Pbf):

10 ^1H NMR (270 MHz, d_6 -acetone): δ , 7.81 (d, 2, J = 7.7 Hz), 7.66 (d, 2, J = 7.3 Hz), 7.29-7.38 (dt, 4), 6.54 (s, br, 1), 4.31 (m, 2), 4.18 (t, 1, J = 6.95 Hz), 4.06 (q, 2, J = 6.95 Hz), 3.63 (m, 1), 3.53 (m, 2), 3.21 (m, 2), 2.94 (m, 2), 2.60 (s, 4), 2.51 (s, 3), 1.96 (s, 2), 1.38 (m, 10), 1.19 (t, 2, J = 7 Hz), 0.89 (m, 1).

15 ^{13}C NMR (67.8 MHz, d_6 -acetone): δ , 158.9, 157.3, 145.1 145.0, 138.7, 135.3, 132.8, 128.4, 127.9, 126.1, 125.3, 120.7, 117.5, 86.9, 66.8, 62.3, 60.5, 53.7, 48.1, 43.6, 41.6, 20.8, 19.5, 18.2, 14.5, 12.5.

Asp(OtBu):-84%

20 ^1H NMR (270 MHz, d_6 -acetone): δ , 7.88 (d, 2, J = 7.5 Hz), 7.69 (d, 2, J = 7.3 Hz), 7.30-7.45 (dt, 4), 7.16 (d, 1, J = 8.8 Hz), 4.84 (m, 1), 4.31 (m, 3), 3.87 (m, 1), 3.39 (m, 2), 2.26 (m, 1), 1.37 (s, 9).

25 ^{13}C NMR (67.8 MHz, d_6 -acetone): δ , 170.4, 155.5, 143.8, 140.7, 127.5, 127.0, 125.1, 120.0, 79.6, 65.2, 62.9, 50.2, 46.7, 27.6.

Asn:-56%

^1H NMR (270 MHz, d_6 -acetone): δ , 7.96, 7.88 (s, d, 2, J = 7.5 Hz),

7.70, (m, 2), 7.31-7.43 (dt, 4), 4.34 (m, 4), 3.74 (m, 3), 3.62 (t, 1, $J = 4.6$ Hz), 3.44 (m, 1), 2.52 (m, 2).

^{13}C NMR (67.8 MHz, d_6 -acetone): δ , 162.3, 155.6, 143.7, 140.6, 127.6, 125.2, 125.0, 120.0, 65.2, 54.1, 46.6, 46.5.

5 Gln:-86%

^1H NMR (270 MHz, d_6 -acetone): δ , 7.96, 7.87 (s, d, 3, $J = 7.3$ Hz), 7.70 (d, 2, $J = 7.1$ Hz), 7.30-7.42 (dt, 4), 7.13 (d, 1, $J = 8.6$ Hz), 4.74 (s, br, 1), 4.24 (m, 3), 3.43 (m, 5), 2.89 (s, 2), 2.73 (s, 2), 2.45 (m, 3), 2.03 (s, 3), 1.80 (m, 1), 1.59 (m, 1).

10 ^{13}C NMR (67.8 MHz, d_6 -acetone): δ , 162.2, 155.9, 143.9, 143.8, 140.7, 127.5, 126.9, 125.1, 124.7, 120.0, 65.1, 63.2, 52.0, 46.7, 35.7, 30.5, 29.9, 14.6.

Glu(OtBu):-71%

15 ^1H NMR (270 MHz, d_6 -acetone): δ , 7.88 (d, 2, $J = 7.3$ Hz), 7.71 (d, 2, $J = 7.3$ Hz), 7.31-7.45 (dt, 4), 7.05 (d, 1, $J = 8.6$ Hz), 4.24 (m, 3), 4.02 (q, 1, $J = 7.1$ Hz), 3.37 (m, 2), 2.20 (m, 2), 1.40 (s, 9).

20 ^{13}C NMR (67.8 MHz, d_6 -acetone): δ , 172.1, 155.9, 143.9, 143.8, 140.7, 127.5, 126.7, 125.2, 120.0, 79.3, 65.1, 63.3, 52.1, 46.8, 31.5, 27.7, 26.3

Gly:-79%

^1H NMR (270 MHz, d_6 -acetone): δ , 7.88 (d, 2, $J = 7.1$ Hz), 7.69 (d, 2, $J = 7.1$ Hz), 7.24-7.42 (m, 5), 4.66 (t, 1, $J = 5.5$ Hz), 4.29 (m, 3), 3.38 (m, 2), 3.09 (q, 2, $J = 6$ Hz).

25 ^{13}C NMR (67.8 MHz, d_6 -acetone): δ , 156.2, 143.9, 140.7, 127.5, 127.0, 125.1, 120.0, 65.2, 59.8, 46.7, 43.0.

Ile:-72%

^1H NMR (270 MHz, d_6 -acetone): δ , 7.88 (d, 2, $J = 7.1$ Hz), 7.73 (d,

2, $J = 7.3$ Hz), 7.29-7.42 (m, 4), 7.03 (d, 1, $J = 8.4$ Hz), 4.51 (m, 1), 4.23 (m, 3), 3.42 (m, 3), 0.84 (m, 6).

^{13}C NMR (67.8 MHz, d_6 -acetone): δ , 156.1, 144.0, 143.8, 140.7, 127.5, 126.9, 125.2, 120.0, 65.1, 61.1, 57.0, 46.8, 35.3, 24.6, 15.4, 11.3.

Leu: -88%

^1H NMR (270 MHz, d_6 -acetone): δ , 7.88 (d, 2, $J = 7.5$ Hz), 7.71 (d, 2, $J = 7.3$ Hz), 7.30-7.44 (dt, 4), 6.99 (d, 1, $J = 8.8$ Hz), 4.24 (m, 3), 3.54 (m, 1), 3.35 (m, 2), 1.62 (m, 1), 1.30 (m, 2), 0.87 (m, 6).

^{13}C NMR (67.8 MHz, d_6 -acetone): δ , 155.9, 144.0, 143.8, 140.7, 127.5, 126.9, 125.2, 125.2, 120.0, 65.0, 64.1, 50.9, 46.8, 24.2, 23.4, 21.7.

Lys(Boc): -95%

^1H NMR (270 MHz, d_6 -acetone): δ , 7.88 (d, 2, $J = 7.7$ Hz), 7.72 (d, 2, $J = 7.0$ Hz), 7.30-7.42 (dt, 4), 4.23 (m, 3), 4.01 (q, 1, $J = 7.0$ Hz), 2.89 (m, 2), 1.37 (m, 12), 1.17 (t, 2, $J = 7.1$ Hz), 0.89 (d, 1, $J = 6.6$ Hz).

^{13}C NMR (67.8 MHz, d_6 -acetone): δ , 155.9, 155.5, 143.9, 143.9, 143.3, 140.7, 127.7, 127.5, 127.0, 125.1, 120.0, 77.2, 65.1, 63.5, 59.7, 52.8, 46.7, 46.6, 29.5, 28.2, 22.8, 20.7, 18.7, 14.0.

Met: -52%

^1H NMR (270 MHz, d_6 -acetone): δ , 7.91, 7.72 (m, d, 4, $J = 6.6$ Hz), 7.34 (m, 4), 7.08 (d, 1, $J = 8.1$ Hz), 6.75 (m, 1), 4.28 (m, 3), 3.42 (m, 3), 2.89 (s, 2), 2.74 (s, 2), 2.11 (m, 2), 1.81 (m, 1), 1.56 (m, 1).

^{13}C NMR (67.8 MHz, d_6 -acetone): δ , 162.2, 155.9, 143.9, 140.7, 127.6, 127.0, 125.2, 120.0, 119.9, 65.2, 63.3, 52.6, 46.7, 26.8.

Phe:-94%

¹H NMR (270 MHz, d₆-acetone): δ, 7.87 (d, 2, J = 7.3 Hz), 7.65 (m, 2), 7.24-7.41 (m, 10), 4.81 (t, 1, J = 5.5 Hz), 4.16 (m, 3), 3.67 (m, 1), 3.39 (m, 2), 2.87 (dd, 1, J = 4.9, 8.6 Hz), 2.67 (m, 1).

5 ¹³C NMR (67.8 MHz, d₆-acetone): δ, 155.6, 143.8, 140.6, 139.2, 129.1, 128.0, 127.5, 127.0, 125.8, 125.2, 125.1, 120.0, 65.1, 62.9, 54.6, 46.6.

Pro:-73%

10 ¹H NMR (270 MHz, d₆-acetone): δ, 7.88 (d, 2, J = 6.7 Hz), 7.65 (d, 2, J = 7.3 Hz), 7.31-7.43 (dt, 4), 4.74 (m, 1), 4.30 (m, 3), 3.73 (m, 1), 3.27 (m, 3), 1.86 (m, 4), 1.17 (t, 1, J = 7.1).

¹³C NMR (67.8 MHz, d₆-acetone): δ, 154.1, 143.9, 140.8, 127.6, 127.0, 125.0, 120.0, 66.4, 66.2, 61.8, 61.1, 59.7, 58.9, 58.3, 46.8, 46.3, 27.7, 26.9, 23.2, 22.4, 20.7, 14.0.

15 Ser(tBu):-76%

¹H NMR (270 MHz, d₆-acetone): δ, 7.88 (d, 2, J = 7.5 Hz), 7.11 (d, 2, J = 7.3 Hz), 7.30-7.45 (dt, 4), 6.99 (d, 1, J = 8.1 Hz), 4.24, 3.74, 3.30-3.56 (m, m, m, 11), 1.12 (s, 9).

20 ¹³C NMR (67.8 MHz, d₆-acetone): δ, 155.9, 144.0, 143.9, 140.8, 127.6, 127.1, 125.3, 125.3, 120.1, 72.4, 65.3, 60.7, 53.6, 46.8, 27.4, 18.9.

Thr(tBu):

25 ¹H NMR (270 MHz, d₆-acetone): δ, 7.90 (d, 2, J = 7.3 Hz), 7.70 (d, 2, J = 7.3 Hz), 7.28-7.44 (dt, 4), 6.89 (d, 1, J = 8.1 Hz), 4.23 (m, 3), 3.68 (m, 1), 3.29 (m, 1), 2.73 (m, 1), 1.15 (m, 12).

¹³C NMR (67.8 MHz, d₆-acetone): δ, 155.6, 143.9, 143.7, 140.8, 127.5, 127.1, 125.2, 125.2, 120.2, 72.3, 65.3, 60.7, 53.6, 46.8, 27.3, 18.6, 17.9.

Trp: -89%

¹H NMR (270 MHz, d₆-acetone): δ, 10.81 (s, 1), 7.86 (d, 2, J = 7.3 Hz), 7.67 (m, 3), 7.34 (m, 5), 6.96-7.14 (m, 4), 4.76 (t, 1, J = 5.3 Hz), 4.24 (m, 3), 3.77 (m, 1), 3.41 (m, 2), 2.94, 2.5 (m, 2).

¹³C NMR (67.8 MHz, d₆-acetone): δ, 155.8, 143.9, 140.6, 136.1, 127.5, 127.5, 127.0, 125.2, 125.2, 123.1, 120.7, 112.0, 118.4, 118.1, 111.4, 111.2, 65.2, 62.8, 53.8, 46.7, 26.7.

Tyr (tBu): -88%

¹H NMR (270 MHz, d₆-acetone): δ, 7.87 (d, 2, J = 7.3 Hz), 7.66 (d, 2, J = 7.3 Hz), 7.31-7.41 (m, 4), 7.11 (m, 3), 6.82 (d, 2, J = 8.2 Hz), 4.79, (t, 1, J = 5.5 Hz), 4.13 (m, 3), 3.64 (m, 1), 3.38 (m, 1), 2.82 (dd, 1, J = 4.8, 9.0 Hz), 1.19 (s, 9).

¹³C NMR (67.8 MHz, d₆-acetone): δ, 155.6, 153.0, 143.9, 143.8, 140.6, 133.8, 129.5, 127.5, 126.9, 125.2, 125.1, 123.3, 112.0, 77.4, 65.2, 63.0, 54.6, 46.6, 28.4.

Val: -96%

¹H NMR (270 MHz, d₆-acetone): δ, 7.88 (d, 2, J = 7.3 Hz), 7.73 (d, 2, J = 7.3 Hz), 7.32-7.41 (dt, 4), 4.24 (m, 3), 3.89 (d, 1, J = 6.2 Hz), 3.39 (m, 2), 0.86 (m, 6), 0.54 (t, 1, J = 7.1 Hz).

¹³C NMR (67.8 MHz, d₆-acetone): δ, 156.3, 152.7, 143.9, 143.9, 140.7, 127.5, 126.9, 125.2, 120.0, 65.2, 61.4, 57.9, 46.8, 46.2, 28.4, 19.5, 18.6, 17.9.

APPENDIX 2

Data for Amino Alkyl Nitrophenyl carbonates incl yield (%), NMR
(d/ppm, d₆-acetone).

Ala:,

5 ¹H NMR (270 MHz, d₆-acetone): δ, 8.32 (d, 2, J = 9.1 Hz), 7.85
(d, 2, J = 7.3 Hz), 7.71 (d, 2, J = 7.3 Hz), 7.56 (d, 2, J =
9.3 Hz), 7.30-7.41 (m, 7), 4.07-4.44 (m, 6), 1.24 (d, 3, J =
6.6 Hz).

¹³C NMR (67.8 MHz, d₆-acetone): δ, 206.4, 156.7, 156.5, 153.1,
10 146.1, 145.0, 144.9, 141.9, 128.3, 127.8, 126.0, 125.9, 123.1,
120.7, 72.1, 66.5, 47.7, 46.3, 17.1.

Arg(Pbf):

¹H NMR (270 MHz, d₆-acetone): δ, 8.13 (d, 2, J = 4.9 Hz), 7.68
(d, 4, J = 7.7 Hz), 7.52 (d, 4, J = 7.8 Hz), 7.24-7.37 (d, 10, J =
15 9.3 Hz), 4.07-4.22 (m, 6), 3.87 (m, 9, 1), 3.37 (m, 2), 2.47
(d, 6, J = 3.1 Hz), 1.26 (m, 12).

¹³C NMR (67.8 MHz, d₆-acetone): δ, 207.3, 160, 158.3, 157.7,
154.3, 147.4, 146.2, 146.1, 145.9, 143.1, 139.8, 136.5, 133.9,
129.5, 129.5, 128.9, 127.0, 124.1, 121.8, 121.8, 88.0, 80.2,
20 72.5, 67.9, 67.8, 66.1, 49.2, 49.1, 44.6, 29.7, 27.8, 20.5,
19.3, 19.2, 13.6.

Asp(OtBu): -74%

¹H NMR (270 MHz, d₆-acetone): δ, 8.31 (d, 2, J = 9.5 Hz), 7.86
(d, 2, J = 7.7 Hz), 7.54 (d, 2, J = 9.1 Hz), 7.39 (m, 6), 4.37
25 (m, 4), 4.24 (m, 1), 4.06 (q, 1, J = 6.9 Hz), 2.66 (m, 1),
1.46 (s, 9).

¹³C NMR (67.8 MHz, d₆-acetone): δ, 206.1, 170.3, 156.7, 153.2,

146.4, 145.0, 142.1, 128.5, 126.8, 126.0, 123.1, 120.8, 81.3, 70.7, 67.1, 48.2, 37.9, 28.2.

Asn:10%

5 ^1H NMR (270 MHz, d_6 -acetone): δ , 8.31 (d, 1, J = 9.5 Hz), 8.14 (m, 2), 7.98 (m, 1), 7.83 (d, 1, J = 7.3 Hz), 7.67 (t, 1, J = 8.1 Hz), 7.53 (d, 1, J = 9.2 Hz), 73.0-7.39 (dt, 3), 6.99 (m, 2), 4.41 (m, 2), 4.24 (m, 1), 2.94 (m, 2), 2.79 (s, 2).

10 ^{13}C NMR (67.8 MHz, d_6 -acetone): δ , 206.3, 164.4, 163.1, 156.8, 156.6, 153.1, 144.9, 142.1, 141.6, 128.5, 127.9, 127.8, 126.1, 123.1, 120.8, 116.5, 79.2, 73.8, 69.6, 67.3, 48.0, 36.3, 20.6.

Gln:-17%

15 ^1H NMR (270 MHz, d_6 -acetone): δ , 8.31 (dd, 2, J = 2.7 Hz, 5.5 Hz), 8.16 9dd, 1, J = 2.2 Hz, 4.8 Hz), 7.83 (d, 2, J = 7.3 Hz), 7.68 (m, 2), 7.27-7.54 (dt, m, 6), 7.03 (dd, 1, J = 2.2 Hz, 4.8 Hz), 6.65 (d, 1, J = 8.4 Hz), 4.21-4.46 (m, 5), 2.59 (m, 2), 1.92 (m, 2).

^{13}C NMR (67.8 MHz, d_6 -acetone): δ , 206.2, 157.1, 156.7, 153.3, 145.1, 144.9, 142.1, 128.5, 127.9, 126.8, 126.3, 123.1, 120.8, 116.6, 71.3, 66.9, 50.2, 48.1, 15.3.

20 Glu(OtBu):-76%

^1H NMR (270 MHz, d_6 -acetone): δ , 8.30 (d, 2, J = 9.1 Hz), 7.83 (d, 2, J = 7.3 Hz), 7.69 (m, 2), 7.50 (d, 1, J = 6.9 Hz), 7.39 (m, 4), 4.39 (m, 2), 4.26 (m, 2), 4.06 (m, 1), 2.38 (m, 1), 1.44 (s, 9).

25 ^{13}C NMR (67.8 MHz, d_6 -acetone): δ , 206.2, 172.6, 157.1, 156.7, 153.2, 146.3, 145.1, 144.9, 142.1, 128.5, 127.9, 126.0, 123.1, 120.8, 80.5, 71.4, 66.9, 50.4, 49.0, 32.3, 27.0.

Gly:-30%

^1H NMR (270 MHz, d_6 -acetone): δ , 8.29 (d, 2, J = 9.1 Hz), 7.84 (d, 2, J = 7.3 Hz), 7.68, (d, 2, J = 7.3 Hz), 7.54, (d, 2, J = 9.1 Hz), 7.31-7.4, (dt, 4), 4.39 (m, 4), 4.24, (t, 1, J = 6.9 Hz), 3.56, (q, 2, J = 5.1 Hz).

5 ^{13}C NMR (67.8 MHz, d_6 -acetone): δ , 206.1, 157.4, 156.7, 153.3, 146.4, 145.1, 142.1, 128.5, 127.9, 126.3, 126.0, 123.1, 120.8, 68.8, 67.1, 48.1, 40.4.

Ile: -91%

10 ^1H NMR (270 MHz, d_6 -acetone): δ , 8.26 (d, 2, J = 9.2 Hz), 7.82 (d, 2, J = 7.5 Hz), 7.68 (m, 2), 7.50 (d, 2, J = 9.1 Hz), 7.26-7.47 (dt, 3), 4.20-4.51 (m, 3), 3.92 (m, 1), 1.60 (m, 1), 1.19 (m, 1), 1.01, 0.92 (d, t, 3, J = 6.8 Hz, 7.3 Hz).

15 ^{13}C NMR (67.8 MHz, d_6 -acetone): δ , 206.3, 156.7, 153.3, 145.4, 144.1, 142.1, 128.5, 127.9, 127.8, 126.1, 126.0, 123.1, 120.8, 70.2, 66.9, 54.9, 54.8, 48.1, 36.7, 15.7, 11.4.

Leu: -88%

20 ^1H NMR (270 MHz, d_6 -acetone): δ , 8.27 (d, 2, J = 9.1 Hz), 7.83 (d, 2, J = 7.3 Hz), 7.67 (t, 2, J = 5.8 Hz), 7.51 (m, 2), 7.29-7.49 (dt, 4), 4.38 (m, 3), 4.22 (m, 3), 1.77 (m, 1), 1.55 (m, 1), 1.37 (m, 1), 0.94 (m, 6).

^{13}C NMR (67.8 MHz, d_6 -acetone): δ , 205.4, 156.4, 155.9, 152.5, 145.6, 144.4, 144.2, 141.3, 127.7, 127.1, 127.1, 125.2, 122.3, 112.0, 71.3, 66.1, 48.2, 47.3, 24.5, 22.7, 21.3.

Lys(Boc): -64%

25 ^1H NMR (270 MHz, d_6 -acetone): δ , 8.30 (m, 1), 7.86 (m, 2), 7.52 (m, 2), 7.31-7.40 (m, 5), 4.23-4.37 (m, 3), 3.08 (m, 2), 1.39 (m, 14).

^{13}C NMR (67.8 MHz, d_6 -acetone): δ , 206.2, 157.2, 156.7, 146.4,

145.1, 145.0, 142.1, 128.5, 127.9, 126.8, 126.0, 123.7, 123.1, 120.8, 116.5, 78.4, 71.6, 67.3, 66.9, 50.9, 48.1, 48.0, 40.8, 40.7, 23.8.

Met: -13%

5 ^1H NMR (270 MHz, d_6 -acetone): δ , 8.29 (d, 2, J = 9.1 Hz), 7.83 (d, 2, J = 7.3 Hz), 7.69 (d, 2, J = 7.3 Hz), 7.39 (m, 4), 4.17-4.42 (m, 3), 3.60-3.71 (m, 2).

^{13}C NMR (67.8 MHz, d_6 -acetone): δ , 205.5, 156.4, 144.4, 144.2, 141.3, 127.7, 127.7, 125.3, 125.3, 122.3, 120.0, 70.6, 66.0, 50.8, 47.4, 29.3, 29.1, 13.5.

Phe :- 36.%

15 ^1H NMR (270 MHz, d_6 -acetone): δ , 8.28 (d, 2, J = 9.1 Hz), 7.82 (d, 2, J = 7.5 Hz), 7.62 (m, 2), 7.51, 7.38-7.18 (d, m, m, 10, J = 9.1 Hz), 6.75 (d, 1, J = 7.9 Hz), 4.46 (d, 1, J = 6.59), 4.31 (m, 3), 4.18 (m, 1), 2.97 (m, 2).

^{13}C NMR (67.8 MHz, d_6 -acetone): δ , 206.2, 156.9, 156.7, 153.2, 146.4, 145.0, 142.0, 138.8, 130.1, 129.2, 128.4, 127.9, 127.3, 126.8, 126.0, 123.1, 120.7, 70.9, 66.9, 52.4, 48.0, 37.7.

Pro: -50.%

20 ^1H NMR (270 MHz, d_6 -acetone): δ , 8.30 (d, 2, J = 8.8 Hz), 7.84, (d, 2, J = 7.7 Hz), 7.67 (d, 2, J = 7.7 Hz), 7.54 (d, 2, J = 9.1 Hz), 7.31-7.40 (m, 4), 4.37, 4.29, 4.04 (m, 6), 3.45 (m, 2), 2.05 (m, 1), 1.96 (m, 3).

25 ^{13}C NMR (67.8 MHz, d_6 -acetone): δ , 205.4, 155.9, 154.8, 152.5, 145.6, 144.4, 141.4, 127.7, 127.2, 125.2, 125.0, 122.4, 120.0, 69.0, 66.8, 55.9, 47.4, 27.4, 23.7, 20.0.

Ser(tBu) :-53%

^1H NMR (270 MHz, d_6 -acetone): δ , 8.31 (m, 1), 7.84 (d, 2, J =

7.7 Hz), 7.69 (d, 2, $J = 7.5$ Hz), 7.53 (d, 1, $J = 9.4$ Hz),
7.42-7.31 (m, 5), 4.35 (m, 4), 3.65 (m, 4), 2.93 (d, 1, $J =$
3.3 Hz), 1.17 (m, 11), 0.92 (d, 1, $J = 6.6$ Hz).

5 ^{13}C NMR (67.8 MHz, d_6 -acetone): δ , 206.1, 156.9, 156.7, 153.2,
145.1, 142.1, 128.5, 128.4, 127.9, 126.1, 126.0, 120.7, 116.5,
79.2, 73.7, 73.4, 69.2, 67.0, 66.9, 62.4, 61.7, 54.2, 51.3,
48.1, 48.0, 47.9, 27.2, 19.2.

Thr(tBu): -45%

10 ^1H NMR (270 MHz, d_6 -acetone): δ , 8.30 (d, 2, $J = 9.1$ Hz), 7.84
(d, 2, $J = 7.7$ Hz), 7.68 (d, 2, $J = 7.3$ Hz), 7.51 (d, 2, $J =$
9.1 Hz), 7.28-7.49 (dt, 4), 6.4 (d, 1, $J = 9.2$ Hz), 4.46,
4.39, 4.24 (dd, d, m, 5, $J = 4.8$ Hz, 10.9 Hz, 7.3 Hz), 3.96-
4.03 (m, 2), 1.21 (m, 12).

15 ^{13}C NMR (67.8 MHz, d_6 -acetone): δ , 205.4, 156.6, 155.9, 152.5,
145.6, 144.4, 144.2, 141.3, 127.7, 127.1, 125.2, 122.3, 120.0,
73.7, 68.3, 66.3, 65.9, 60.6, 55.4, 54.9, 54.1, 47.3, 18.7.

Trp: -64%

20 ^1H NMR (270 MHz, d_6 -acetone): δ , 8.30 (d, 2, $J = 9.1$ Hz), 7.85
(d, 2, $J = 7.7$ Hz), 7.69 (m, 3), 7.52 (d, 2, $J = 8.8$ Hz),
7.24-7.42 (m, 2), 4.35 9m, 6), 3.08 (d, 2, $J = 4.0$ Hz).

^{13}C NMR (67.8 MHz, d_6 -acetone): δ , 206.4, 156.9, 156.4, 153.0,
146.1, 144.1, 144.8, 141.8, 137.4, 128.3, 127.7, 126.0, 125.7,
124.2, 123.1, 121.7, 120.6, 119.2, 119.0, 112.2, 70.8, 66.5,
51.5, 47.8, 27.5.

25 Tyr(tBu): -50%

^1H NMR (270 MHz, d_6 -acetone): δ , 8.30 (d, 2, $J = 9.1$ Hz), 7.84
(d, 2, $J = 7.7$ Hz), 7.64 (d, 2, $J = 7.3$ Hz), 7.52 (d, 2, $J =$
9.1 Hz), 7.29-7.38 9m, 6), 6.90 (d, 2, $J = 8.42$ Hz), 4.14-4.45

(m, 7), 2.89 (m, 4), 1.24 (s, 10).

^{13}C NMR (67.8 MHz, d_6 -acetone): δ , 206.2, 156.9, 156.7, 155.1, 153.2, 146.3, 145.0, 142.0, 133.3, 130.8, 130.5, 128.4, 127.9, 126.0, 124.7, 123.1, 120.7, 78.4, 70.9, 66.9, 52.5, 52.4,

5 48.0, 37.0.

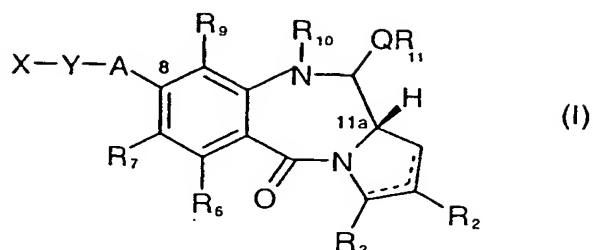
Val :- 62%

^1H NMR (270 MHz, d_6 -acetone): δ , 8.30 (d, 2, J = 9.1 Hz), 7.83 (d, 2, J = 7.3 Hz), 7.70 (d, 2, J = 7.1 Hz), 7.54 (d, 2, J = 9.2 Hz), 7.30-7.44 (m, 4), 4.10-4.44 (m, 6), 3.89 (d, 1, J = 10 6.2 Hz), 0.86 (m, 6).

^{13}C NMR (67.8 MHz, d_6 -acetone): δ NMR (67.8 MHz, d_6 -acetone): δ , 206.2, 156.4, 155.8, 152.3, 146.1, 145.3, 141.9, 128.5, 127.9, 127.1, 125.9, 123.1, 119.9, 71.2, 66.2, 48.1, 47.5, 24.3, 22.7.

CLAIMS

1. A compound of formula (I):



wherein:

5 X is selected from COOH, NHZ, SH, or OH, where Z is either H or an amine protecting group;

A is O, S, NH, or a single bond;

10 R₂ and R₃ are independently selected from: H, R, OH, OR, =O, =CH-R, =CH₂, CH₂-CO₂R, CH₂-CO₂H, CH₂-SO₂R, O-SO₂R, CO₂R, COR, CN and there is optionally a double bond between C1 and C2 or C2 and C3;

R₆, R₇, and R₈ are independently selected from H, R, OH, OR, halo, nitro, amino, Me₃Sn;

R₁₁ is either H or R;

15 Q is S, O or NH;

R₁₀ is a nitrogen protecting group;

20 where R is a lower alkyl group having 1 to 10 carbon atoms, or an aralkyl group, of up to 12 carbon atoms, whereof the alkyl group optionally contains one or more carbon-carbon double or triple bonds, which may form part of a conjugated system, or an aryl group, of up to 12 carbon atoms; and is optionally substituted by one or more halo, hydroxy, amino, or nitro groups, and optionally contains one or more hetero atoms, which may form part of, or be, a functional group; and

Y is a divalent group such that $HY = R$.

5 2. A compound according to claim 1, wherein R and HY are independently selected from lower alkyl group having 1 to 10 carbon atoms, or an aralkyl group, of up to 12 carbon atoms, or an aryl group, of up to 12 carbon atoms, optionally substituted by one or more halo, hydroxy, amino, or nitro groups.

10 3. A compound according to claim 2, wherein R and HY are independently selected from lower alkyl groups having 1 to 10 carbon atoms optionally substituted by one or more halo, hydroxy, amino, or nitro groups.

15 4. A compound according to claim 3, wherein R or HY are independently selected from unsubstituted straight or branched chain alkyl groups, having 1 to 10 carbon atoms.

20 5. A compound according to any one of the preceding claims, wherein R_{10} has a carbamate functionality where it binds to the nitrogen atom at the 10 position of the PBD ring structure.

25 6. A compound according to any one of the preceding claims, wherein R_7 is an electron donating group.

7. A compound according to any one of the preceding claims, wherein Q is O, and/or R_{11} is H.

8. A compound according to any one of the preceding claims,

wherein R_6 and R_9 are H.

9. A compound according to claim 8, wherein R_7 is an alkoxy group.

5

10. A compound according to any one of the preceding claims, wherein R_2 and R_3 are H.

10

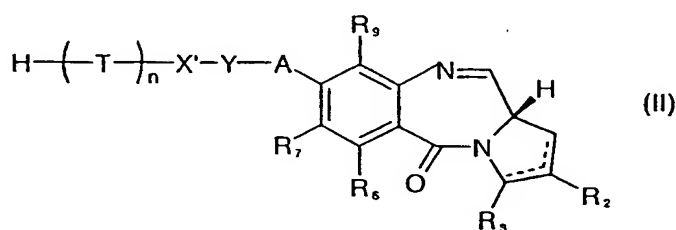
11. A compound according to any one of the preceding claims, wherein there is no double bond between C2 and C3.

12. A compound according to any one of the preceding claims, wherein

-Y-A- is an alkoxy chain.

15

13. A compound of formula II :



wherein Y, A, R_7 , R_2 , R_3 , R_6 , and R_9 are as defined in any one of claims 1 to 12;

X' is CO, NH, S or O;

20

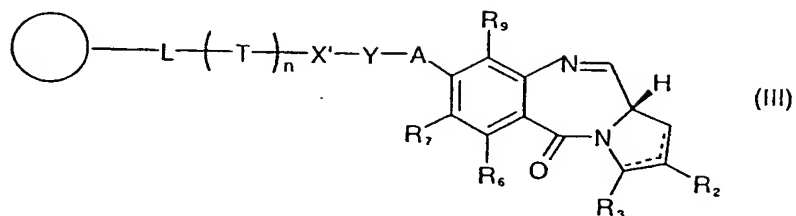
T is a combinatorial unit;

and n is a positive integer.

14. A compound according to claim 13, wherein X' is either CO or NH.

15. A compound according to any one of claims 13 to 15,
wherein n is from 1 to 16.

16. A compound of formula III:



5 wherein X', Y, A, R₇, R₂, R₃, R₆, R₉, and T are as defined in any
one of claims 13 to 15;

n is zero or a positive integer;

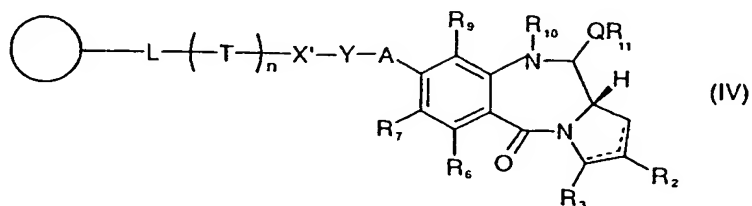
L is a linking group, or a single bond;

and O is a solid support.

10

17. A compound according to claim 18, wherein L is a linking
group.

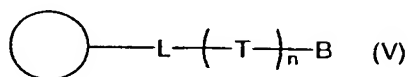
18. A compound of the formula IV:



15

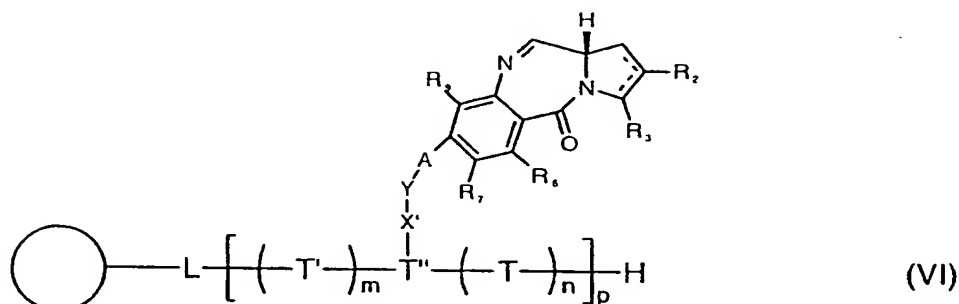
wherein X', Y, A, R₇, R₂, R₃, R₆, R₉, T, n, L and O are as
defined in either claim 16 or claim 17, and R₁₀, R₁₁, and Q are
as defined in any one of claims 1 to 12.

19. A method of making a compound according to claim 18, by reacting a compound of formula I as defined in any of claims 1 to 12 with a compound of formula V:



wherein O, L, T and n are as defined in claim 19, and B is H or an atom or group for providing a functional group capable of reaction with X.

20. A compound of the formula VI:



wherein O, L, X', Y, A, R₂, R₃, R₆, R₇, R₉, and T are as defined in either claim 16 or claim 17;

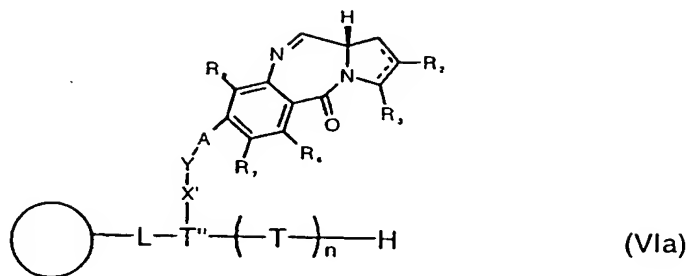
n and m are positive integers, or one of them may be zero;

T' is a combinatorial unit, where each T' may be different if m is greater than 1;

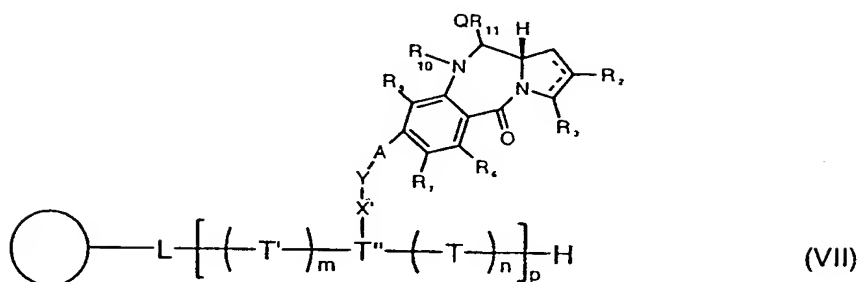
T'' is a combinatorial unit which provides a site for the attachment of X'; and

p is a positive integer, where if p is greater than 1, for each repeating unit, the meaning of X', Y, A, R₂, R₃, R₆, R₇, R₉, T, T', T'' and the values of n and m are independently selected.

21. A compound according to claim 2 of the formula **VIa**:



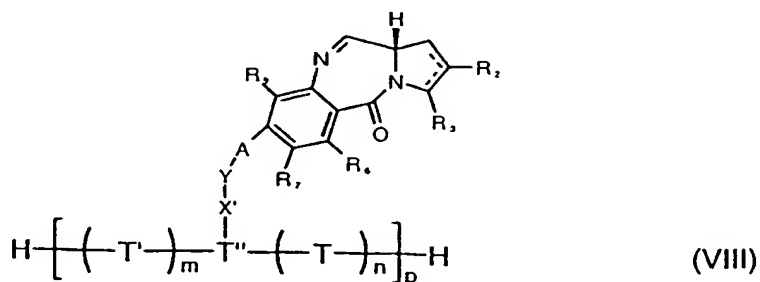
22. A compound of the formula **VII**:



5 wherein O, L, X', Y, A, R₂, R₃, R₆, R₇, R₉, T, T', T'', n, m and p are as defined in claim 20 and Q, R₁₀, and R₁₁, are as defined in claim 18, where if p is greater than 1, for each repeating unit the meanings of X', Y, A, R₂, R₃, R₆, R₇, R₉, T, T', T'', Q, R₁₀, R₁₁ and the values of n and m are independently selected.

10

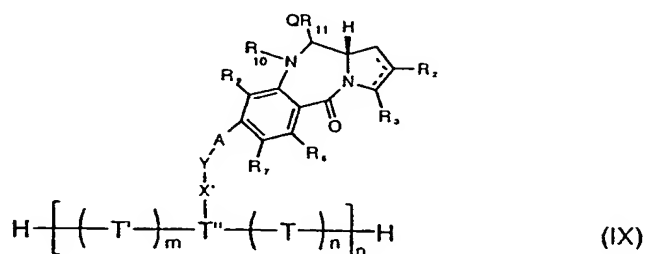
23. A compound of the formula **VIII**:



wherein X' , Y , A , R_2 , R_3 , R_6 , R_7 , R_9 , T , T' , T'' , n , m and p are as defined in the claim 20, where if p is greater than 1, for each repeating unit the meanings of X' , Y , A , R_2 , R_3 , R_6 , R_7 , R_9 , T , T' , T'' and values of n and m are independently selected.

5

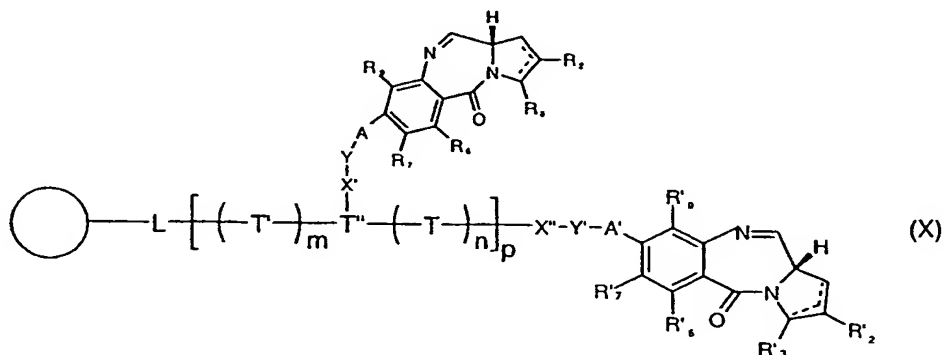
24. A compound of the formula IX:



wherein X' , Y , A , R_2 , R_3 , R_6 , R_7 , R_9 , Q , R_{10} , R_{11} , T , T' , T'' , n , m and p are as defined in claim 22, where if p is greater than 1, for each repeating unit the meanings of X' , Y , A , R_7 , R_2 , R_3 , R_6 , R_9 , T , T' , T'' , Q , R_{10} , R_{11} and values of n and m are independently selected.

10

25. A compound of the formula X:



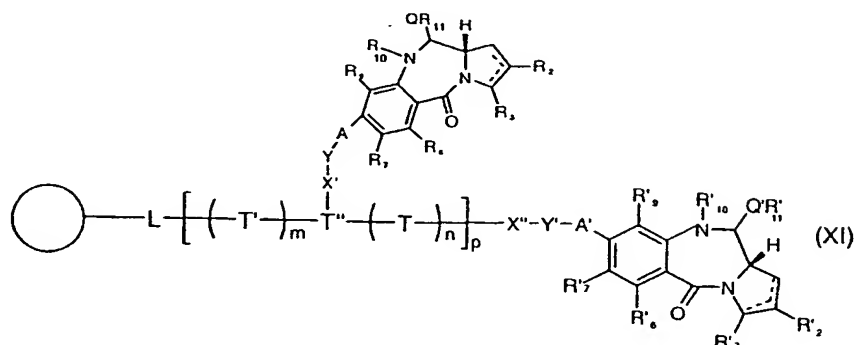
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wherein O , L , X' , Y , A , R_2 , R_3 , R_6 , R_7 , R_9 , T , T' , T'' , n , m and p are as defined in claim 20, and X'' , Y' , A' , R'_2 , R'_3 , R'_6 , R'_7 and R' , are selected from the same possibilities as X' , Y , A ,

R_2 , R_3 , R_6 , R_7 and R_9 , respectively, and where if p is greater than 1, for each repeating unit the meaning of X' , Y , A , R_2 , R_3 , R_6 , R_7 , R_9 , T , T' , T'' and the values of n and m may be independently selected.

5

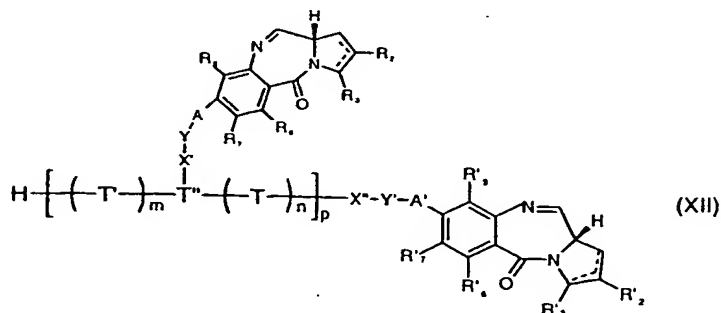
26. A compound of the formula **XI**:



wherein O , L , X' , Y , A , R_2 , R_3 , R_6 , R_7 , R_9 , X'' , Y' , A' , R'_2 , R'_3 , R'_6 , R'_7 , R'_9 , T , T' , T'' , n , m and p are as defined in claim 25, Q , R_{10} , and R_{11} , are as defined in claim 18, and Q' , R'_{10} , R'_{11} , have the same definitions as Q , R_{10} , R_{11} , respectively, and where if p is greater than 1, for each repeating unit the meanings of X' , Y , A , R_2 , R_3 , R_6 , R_7 , R_9 , T , T' , T'' , Q , R_{10} , R_{11} and the values of n and m are independently selected.

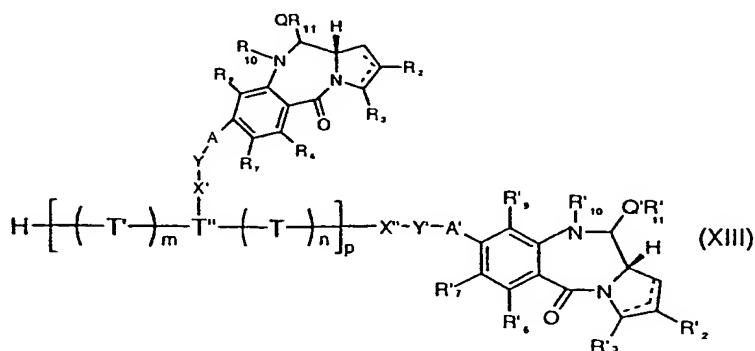
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27. A compound of the formula **XII**:



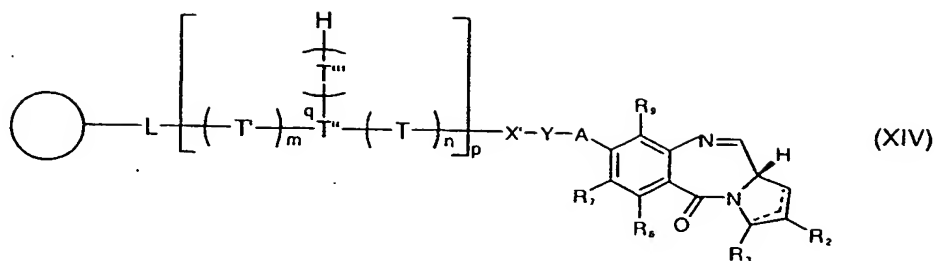
wherein X' , Y , A , R_1 , R_2 , R_3 , R_6 , R_9 , X'' , Y' , A' , R'_1 , R'_2 , R'_3 , R'_6 , R'_9 , T , T' , T'' , n , m and p are as defined in claim 25, and where if p is greater than 1, for each repeating unit the meanings of X' , Y , A , R_1 , R_2 , R_3 , R_6 , R_7 , R_9 , T , T' , and T'' and the values of n and m may be independently selected.

28. A compound of the formula **XIII**:



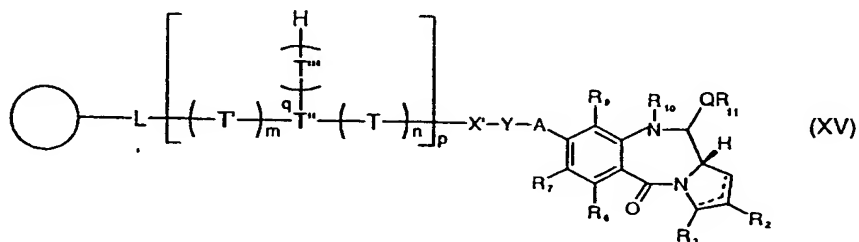
wherein X' , Y , A , R_1 , R_2 , R_3 , R_6 , R_7 , R_9 , Q , R_{10} , R_{11} , X'' , Y' , A' , R'_1 , R'_2 , R'_3 , R'_6 , R'_7 , R'_9 , Q' , R'_{10} , R'_{11} , T , T' , T'' , n , m and p are as defined in claim 26, and where if p is greater than 1, for each repeating unit the meanings of X' , Y , A , R_1 , R_2 , R_3 , R_6 , R_7 , R_9 , T , T' , T'' , Q , R_{10} , R_{11} and the values of n and m may be independently selected.

29. A compound of the formula **XIV**:



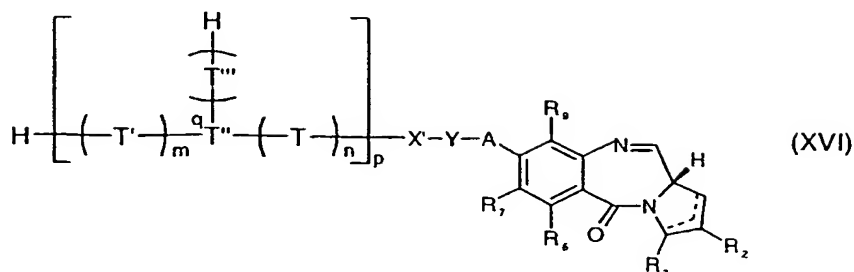
wherein O, L, X', Y, A, R₂, R₃, R₆, R₇, R₉, T, T', T'', n, m and p are as defined in claim 20, and T''' and q are selected from the same possibilities as T and n respectively, and where if p is greater than 1, for each repeating unit the meaning of T, T', T'', T''' and the values of n, m and q may be independently selected.

30. A compound of the formula **XV**:



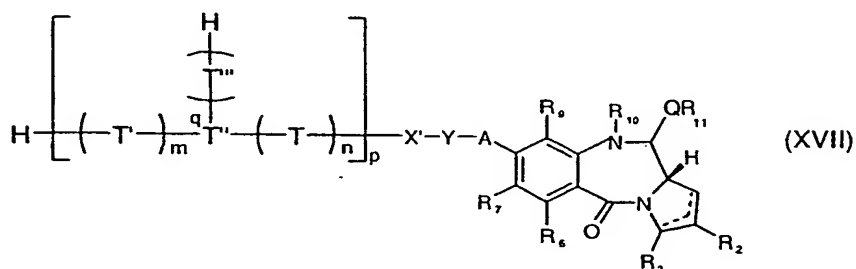
wherein O, L, X', Y, A, R₂, R₃, R₆, R₇, R₉, T, T', T'', T''', n, m, p and q are as defined in claim 29, Q, R₁₀, and R₁₁ are as defined in claim 18, and where if p is greater than 1, for each repeating unit the meanings of T, T', T'', T''' and the values of n, m and q may be independently selected.

31. A compound of the formula **XVI**:



wherein X' , Y , A , R_1 , R_2 , R_3 , R_4 , R_5 , R_6 , R_7 , R_8 , T , T' , T'' , T''' , n , m , p and q are as defined in claim 29, and where if p is greater than 1, the meanings of T , T' , T'' , T''' and values of n , m and q may be independently selected.

32. A compound of the formula **XVII**:



wherein X' , Y , A , R_1 , R_2 , R_3 , R_4 , R_5 , R_6 , R_7 , R_8 , Q , R_{10} , R_{11} , T , T' , T'' , T''' , n , m , p and q are as defined in claim 30, and where if p is greater than 1, for each repeating unit the meanings of T , T' , T'' , T''' and the values of n , m and q may be independently selected.

33. A collection of compounds all of which are represented by either:

- (i) formula I as defined in any one of claims 1 to 12;
- (ii) formula II as defined in any one of claims 13 to 15;
- (iii) formula III as defined in either claim 16 or 17;
- (iv) formula IV as defined in claim 18;
- 5 (v) formula VI as defined in claim 20;
- (vi) formula VII as defined in claim 22;
- (vii) formula VIII as defined in claim 23;
- (viii) formula IX as defined in claim 24;
- (ix) formula X as defined in claim 25;
- 10 (x) formula XI as defined in claim 26;
- (xi) formula XII as defined in claim 27;
- (xii) formula XIII as defined in claim 28;
- (xiii) formula XIV as defined in claim 29;
- (xiv) formula XV as defined in claim 30;
- 15 (xv) formula XVI as defined in claim 31; or
- (xvi) formula XVII as defined in claim 32.

34. A method of preparing a collection of compounds as defined in claim 33.

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35. A method of screening compounds of:

- (i) formula II as defined in any one of claims 13 to 15;
- (ii) formula III as defined in either claim 16 or 17;
- (iii) formula IV as defined in claim 18;
- 25 (iv) formula VI as defined in claim 20;
- (v) formula VIII as defined in claim 23;
- (vi) formula X as defined in claim 25;

- (vii) formula XII as defined in claim 27;
 - (viii) formula XIV as defined in claim 29; or
 - (ix) formula XVI as defined in claim 31
- to discover biologically active compounds.

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36. The use of a compound of:

- (i) formula II as defined in any one of claims 13 to 15;
- (ii) formula VIII as defined in claim 23;
- (iii) formula XII as defined in claim 27; or
- (iv) formula XVI as defined in claim 31

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in the manufacture of a cytotoxic, antibiotic, antiparasitic or antiviral therapeutic composition.

37. The use of a compound of:

- (i) formula II as defined in any one of claims 13 to 15;
- (ii) formula VIII as defined in claim 23;
- (iii) formula XII as defined in claim 27; or
- (iv) formula XVI as defined in claim 31

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in a method of diagnosis.

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38. The use of a compound of:

- (i) formula II as defined in any one of claims 13 to 15;
- (ii) formula VIII as defined in claim 23;
- (iii) formula XII as defined in claim 27; or
- (iv) formula XVI as defined in claim 31

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in a method of target validation pr om functional genomics.

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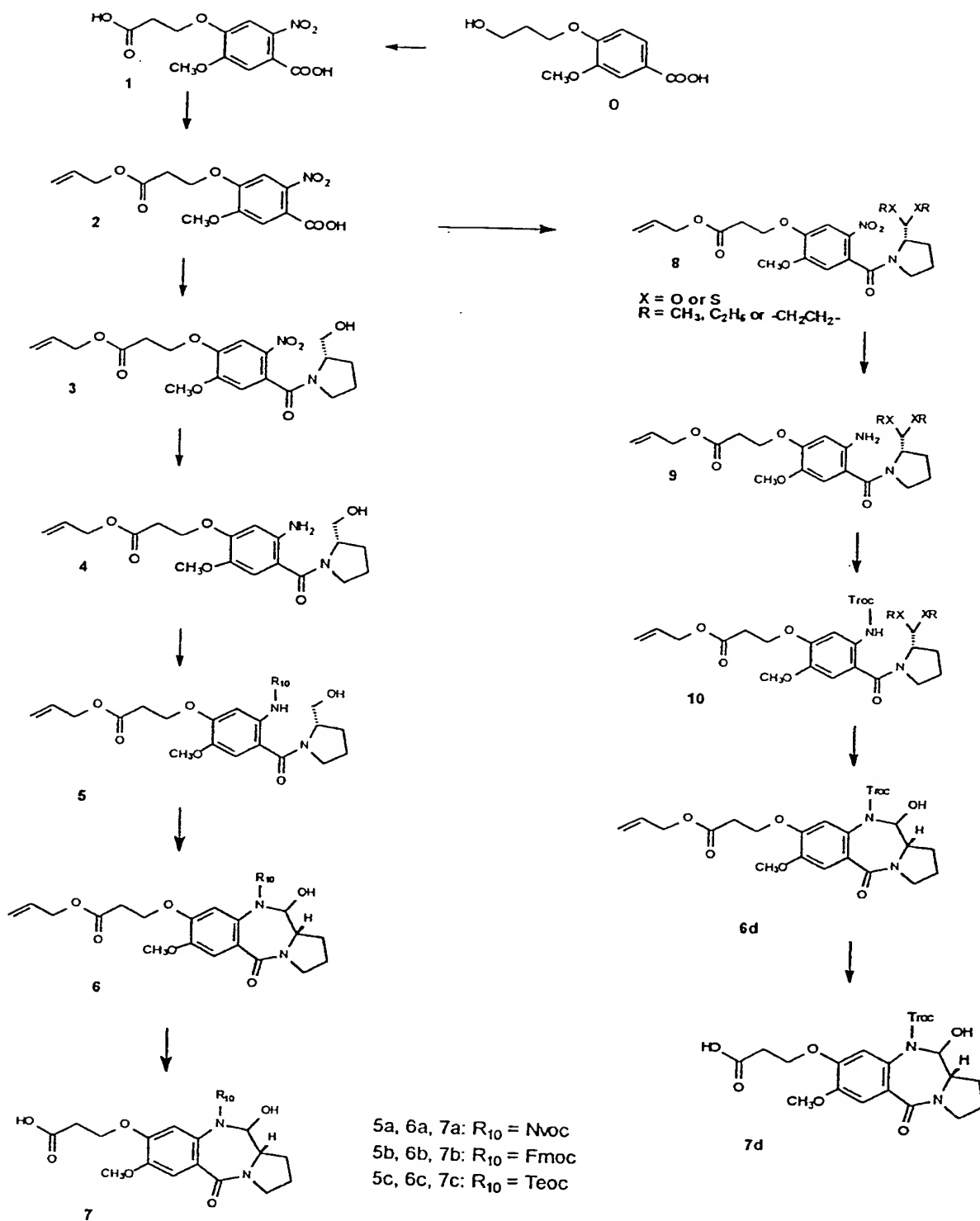


Figure 1

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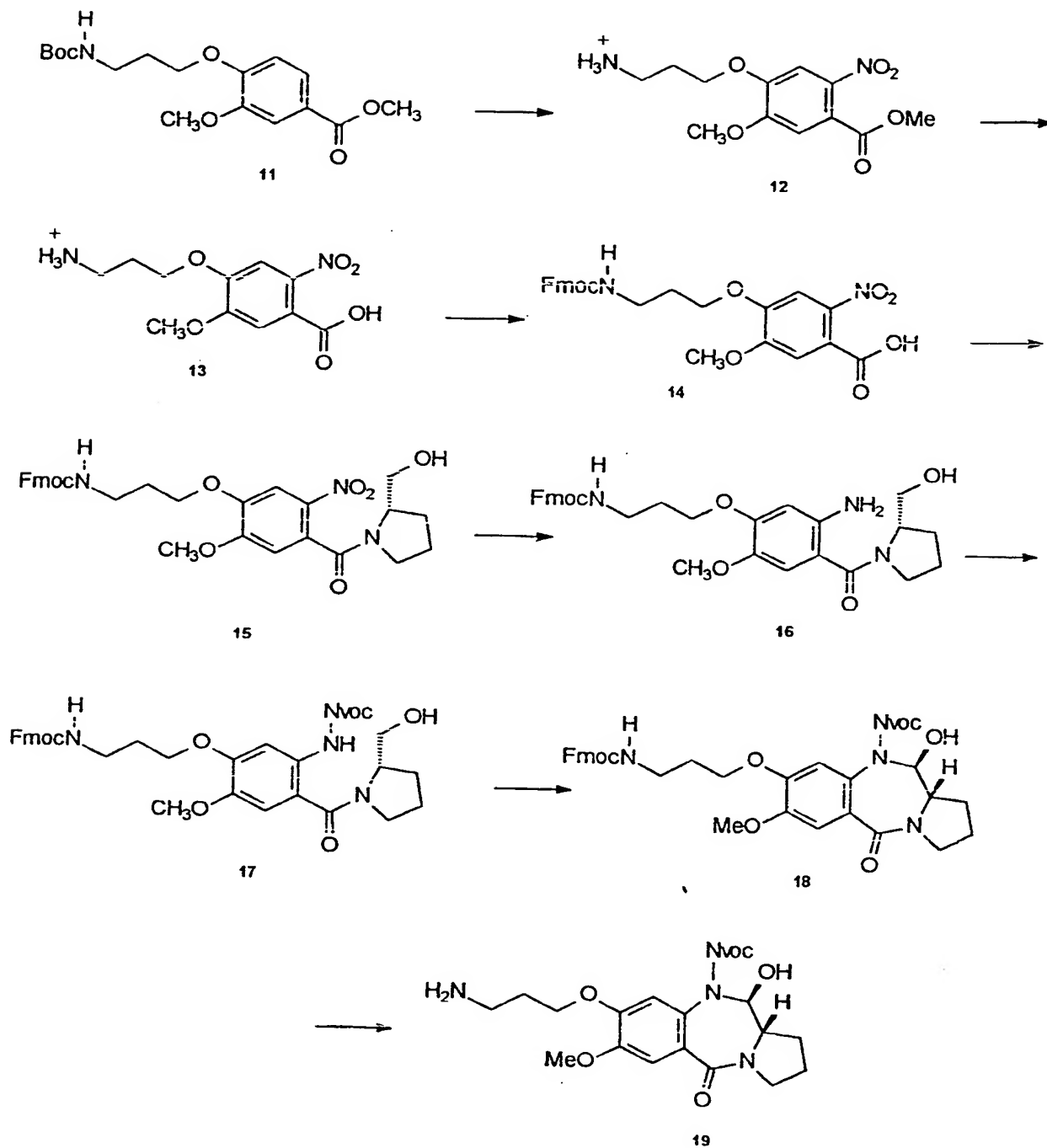
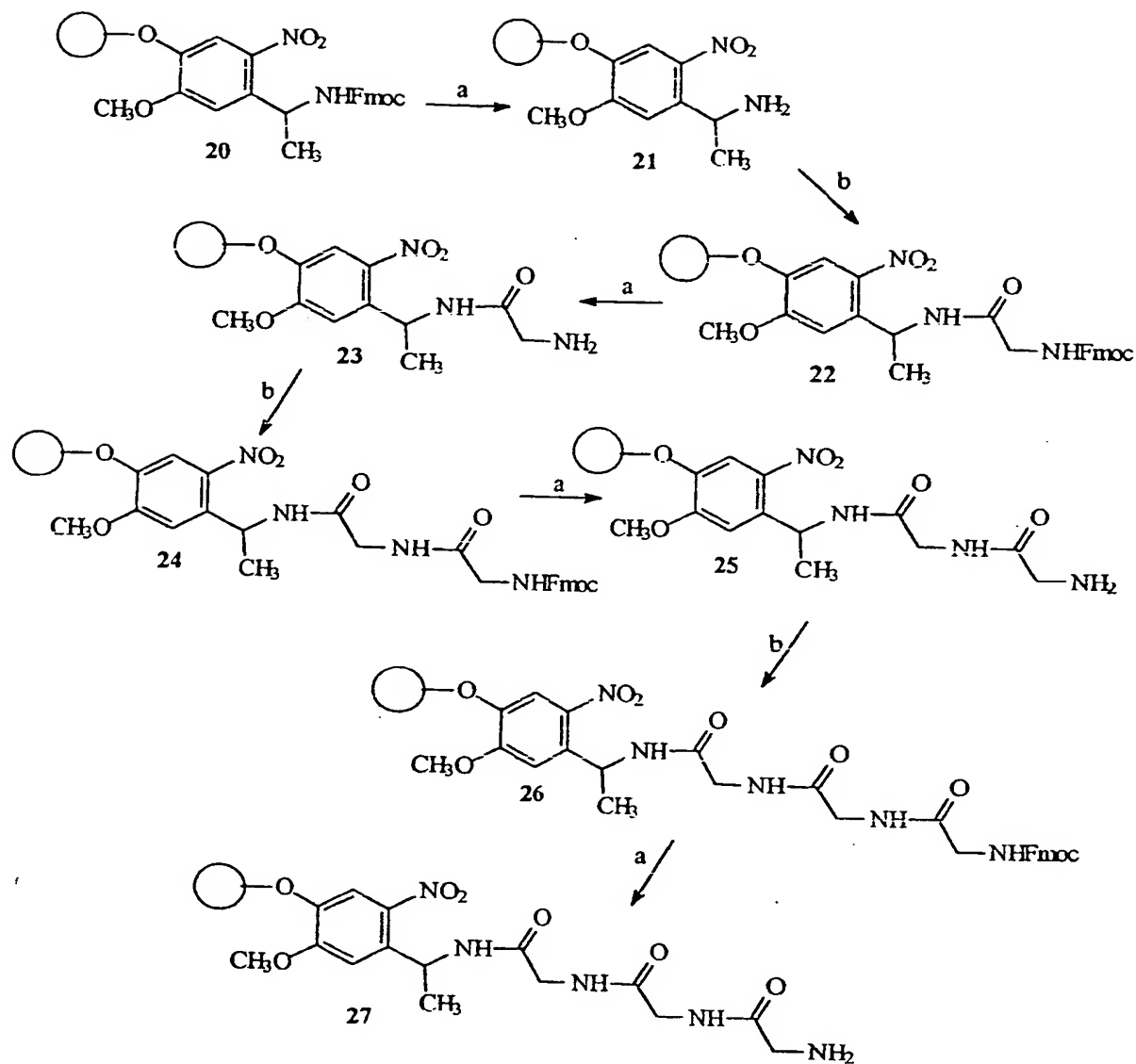


Figure 2

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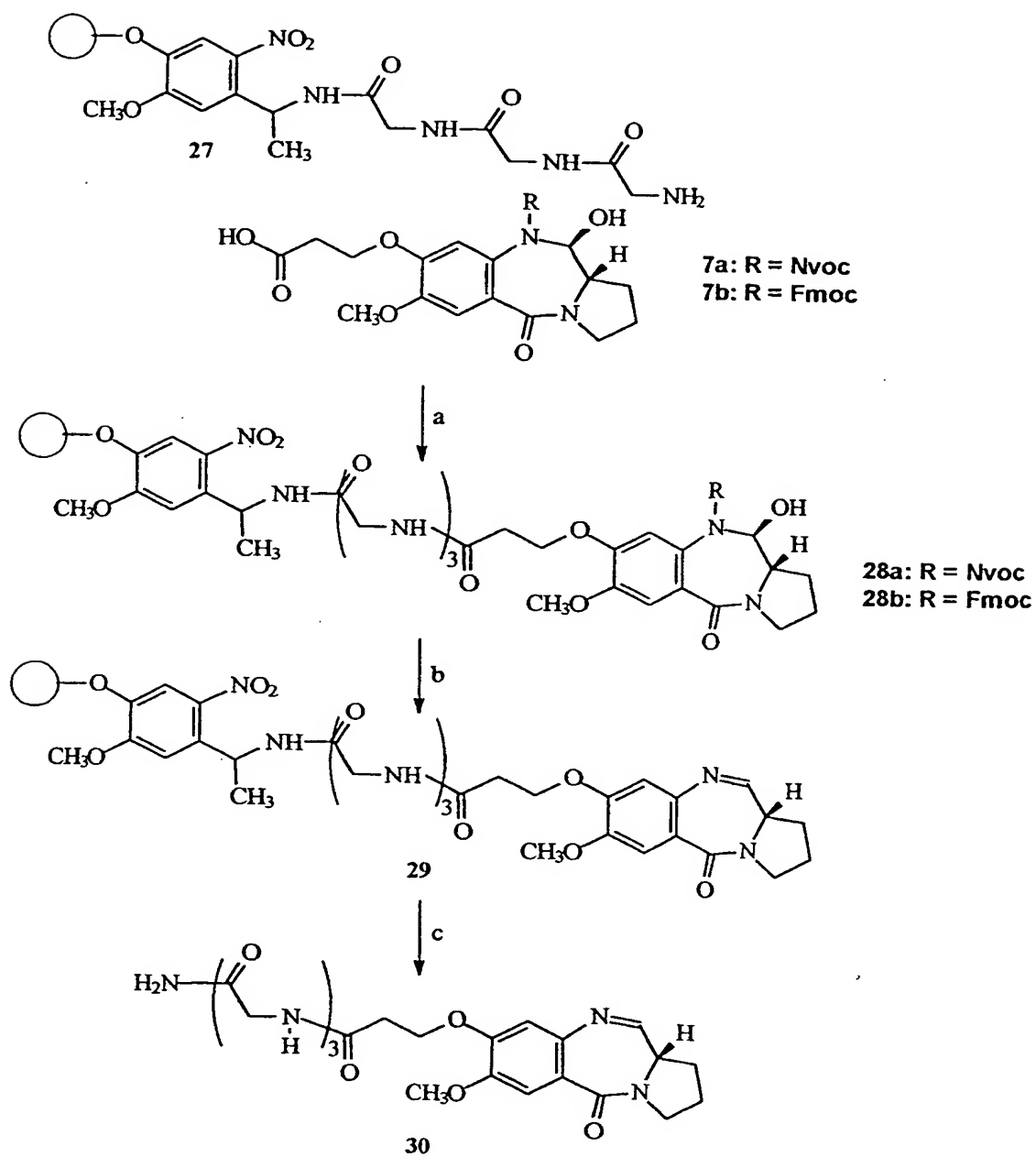


a: 20% piperidine/DMF; b: (i) Fmoc.glycine, TBTU, DIPEA, DMF,
(ii) 20% Ac₂O, 30% pyridine, 50% CH₂Cl₂

Figure 3a

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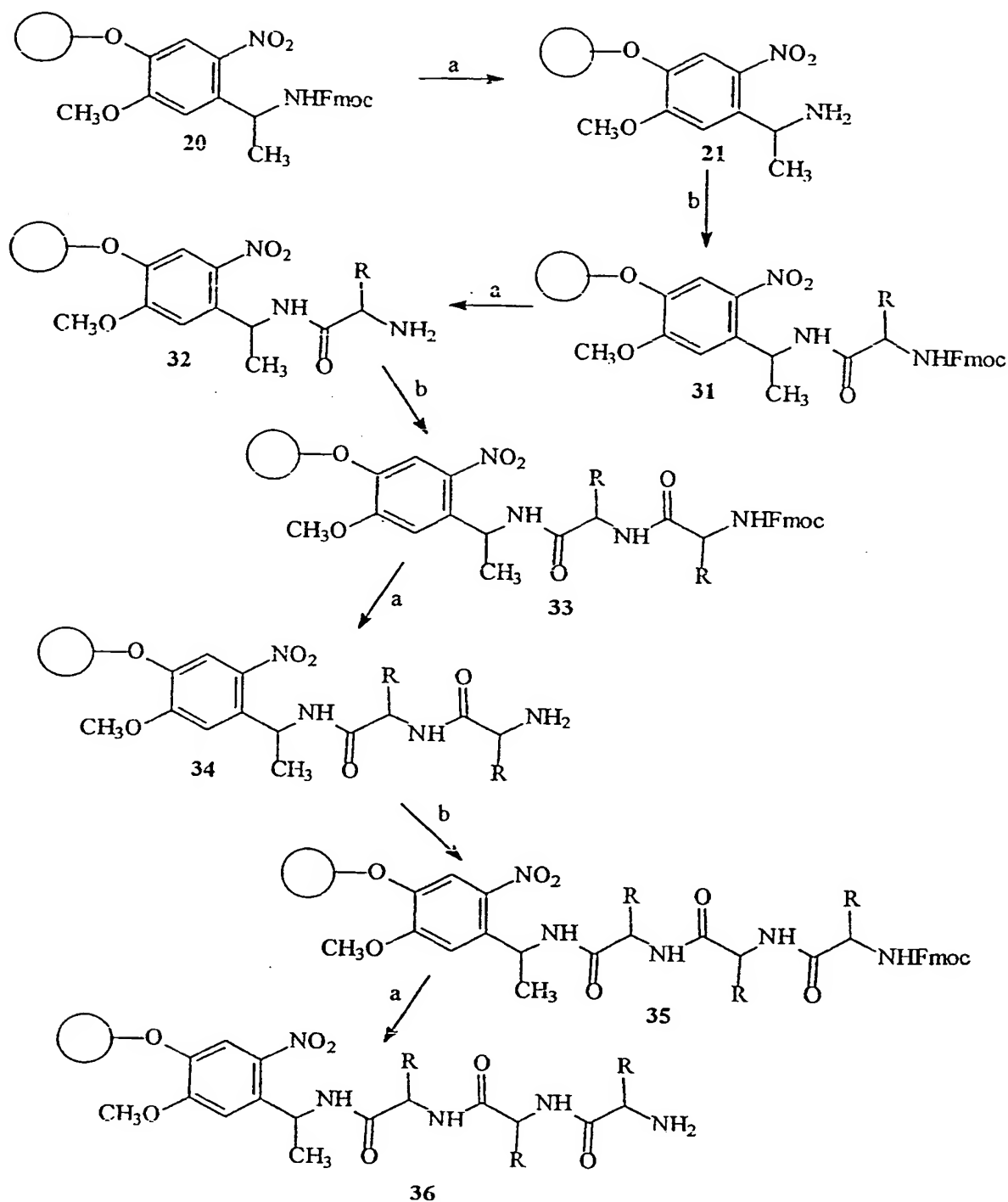


a: (i) TBTU, DIPEA, DMF, (ii) 20% Ac₂O, 30% pyridine, 50% CH₂Cl₂;
 b: 20% piperidine/DMF; c: hn

Figure 3b

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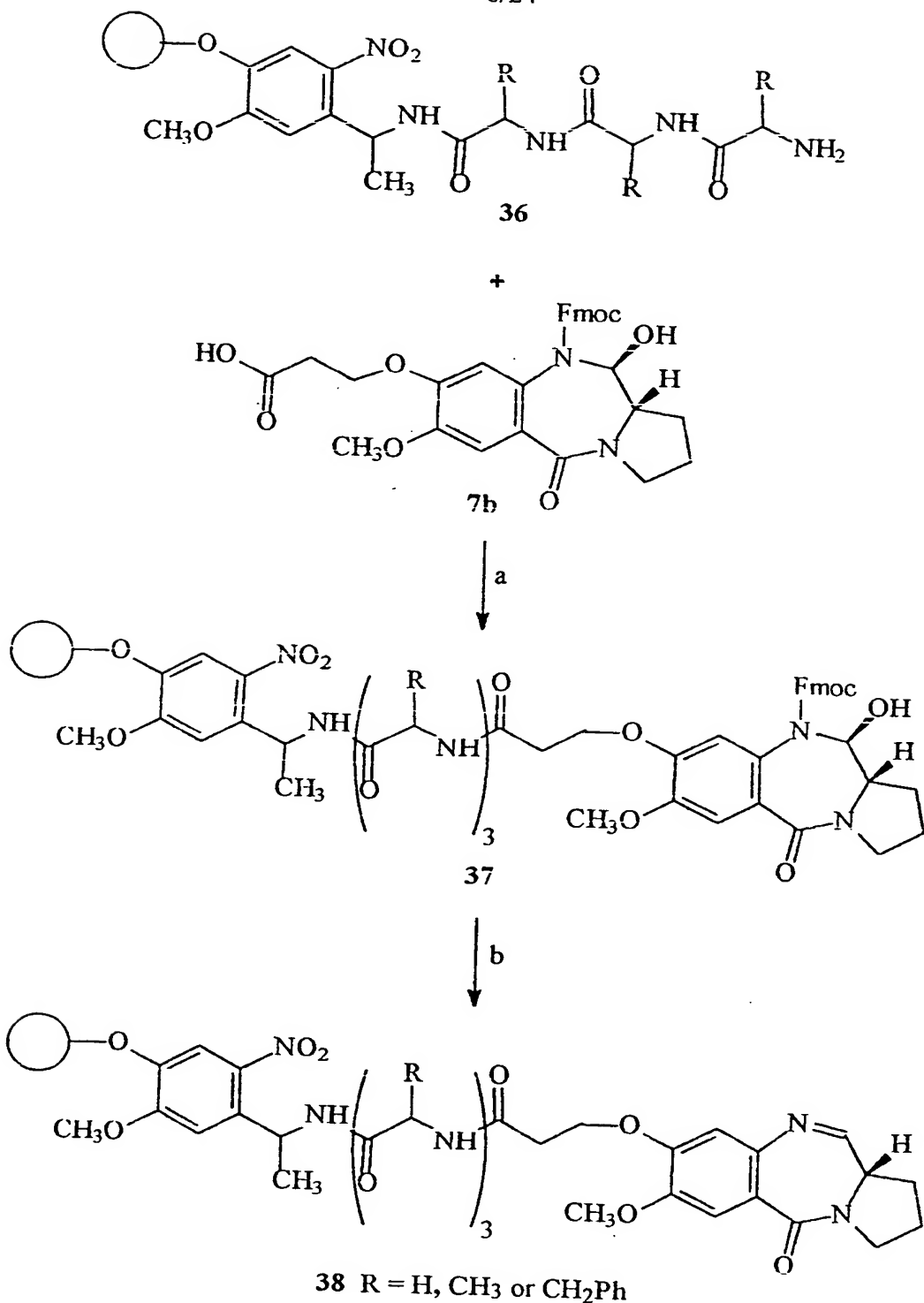


a: 20% piperidine/DMF; b: (i) Fmoc.glycine, valine, phenylalanine, TBTU, DIPEA, DMF;
(ii) 20% Ac_2O , 30% pyridine, 50% CH_2Cl_2

Figure 4a

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a: (i) TBTU, DIPEA, DMF; (ii) 20% Ac₂O, 30% pyridine, 50% CH₂Cl₂;
 b: 20% piperidine/DMF

Figure 4b

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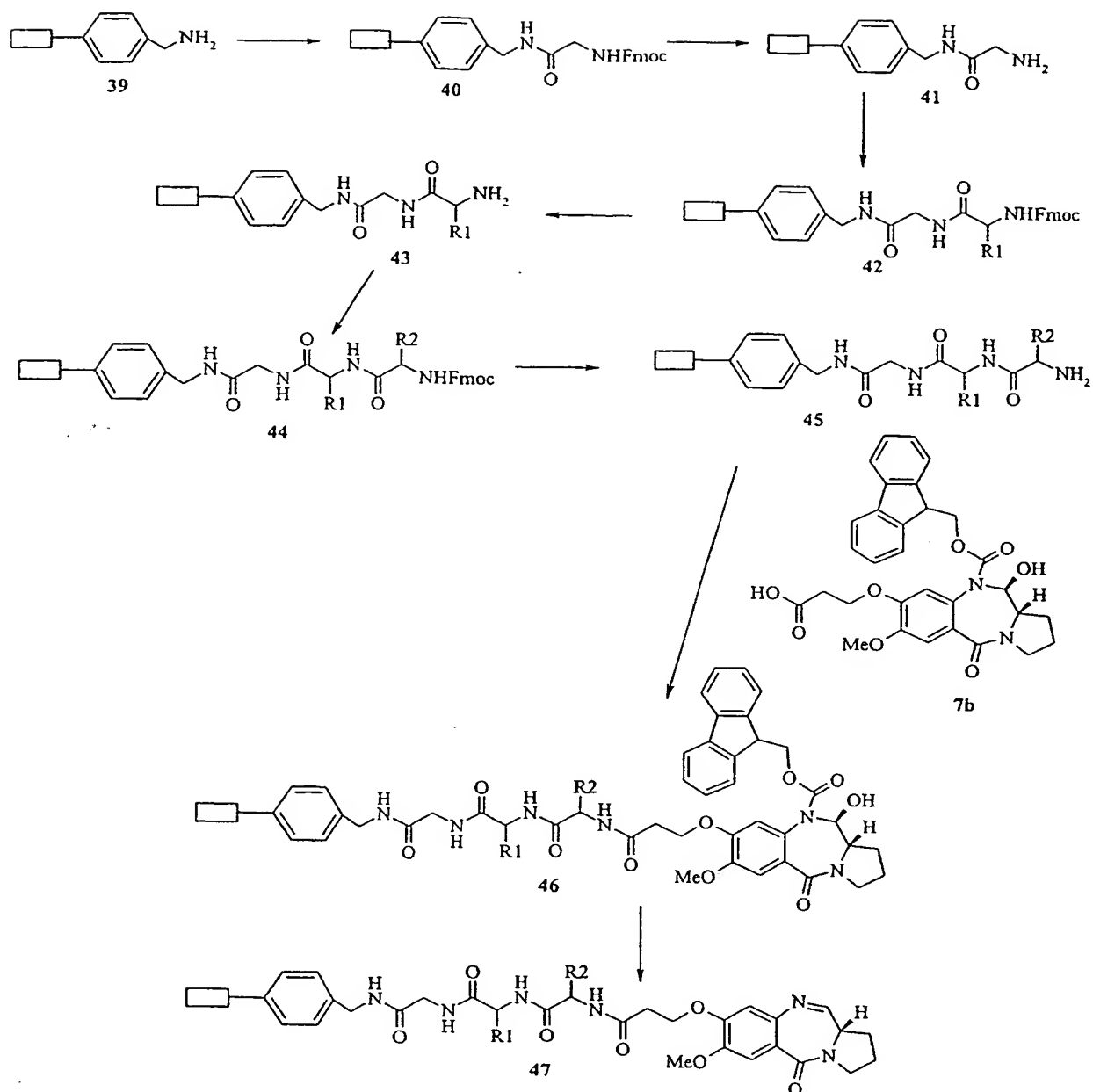


Figure 5

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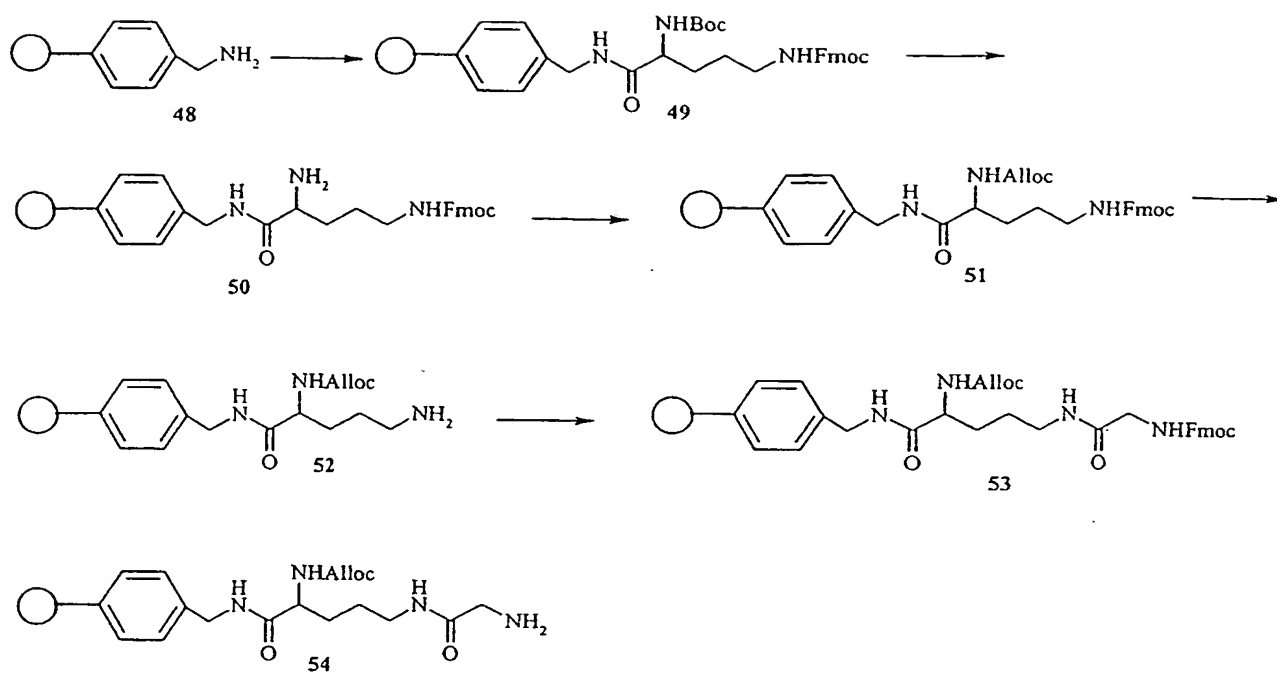


Figure 6a

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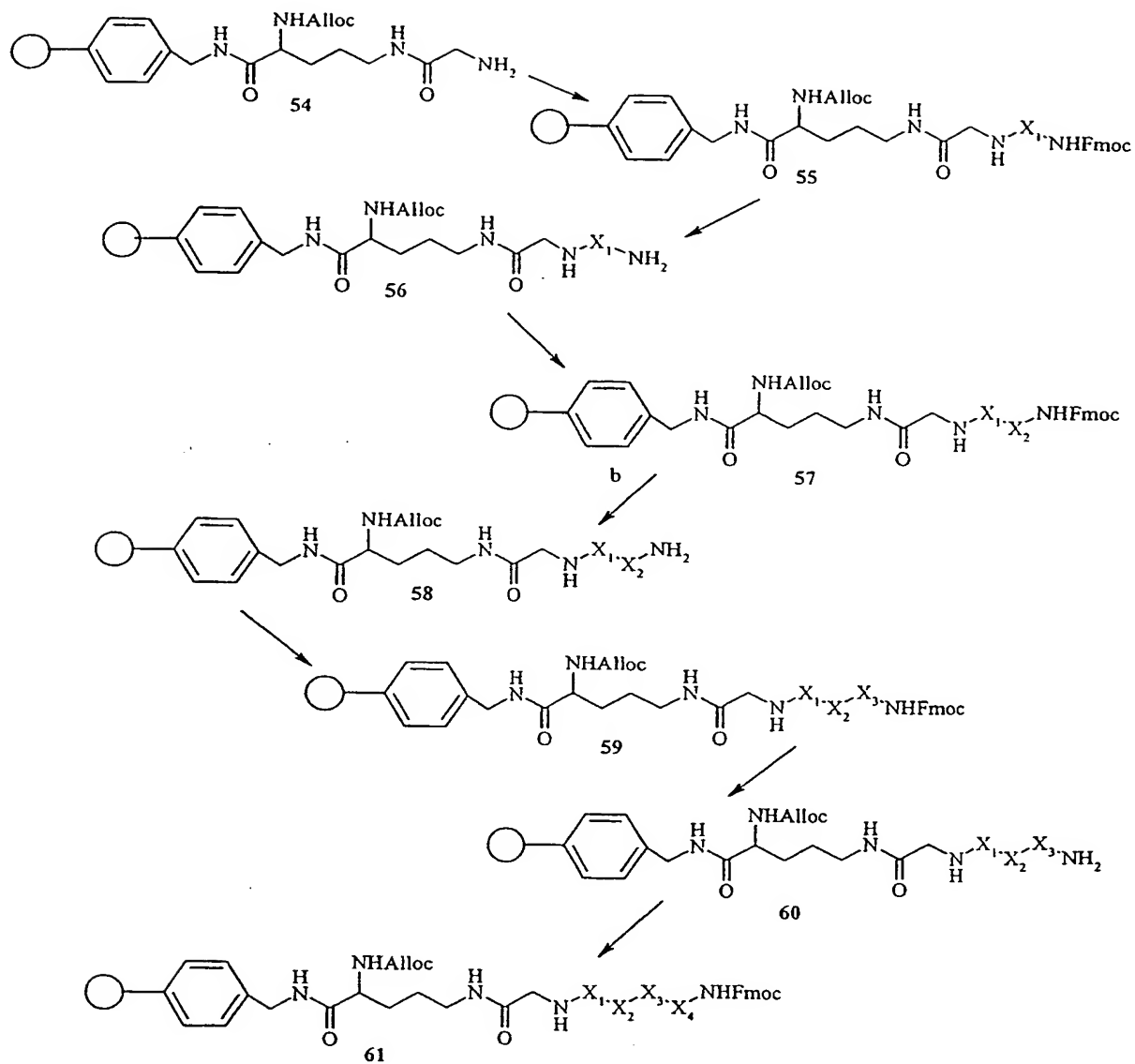


Figure 6b

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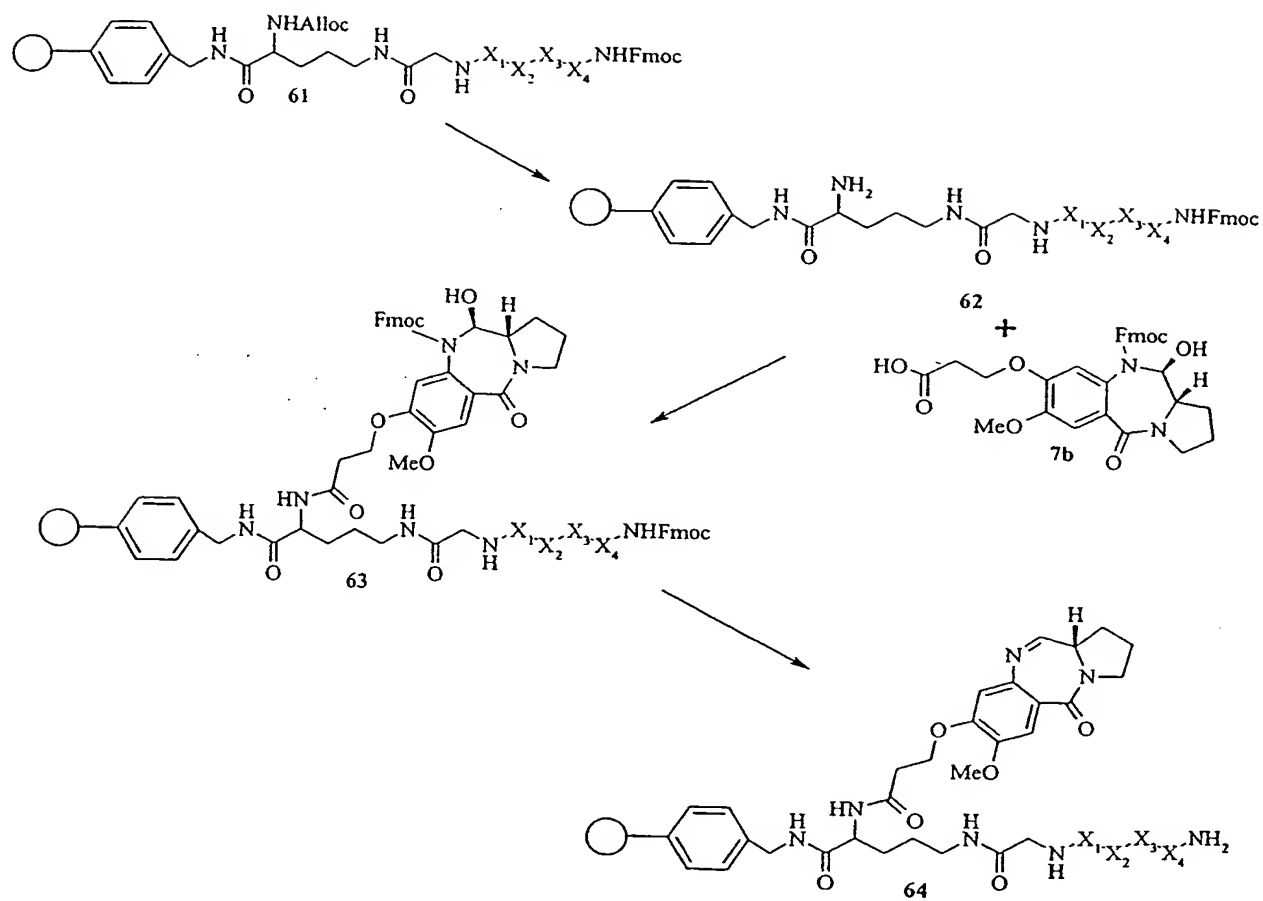


Figure 6c

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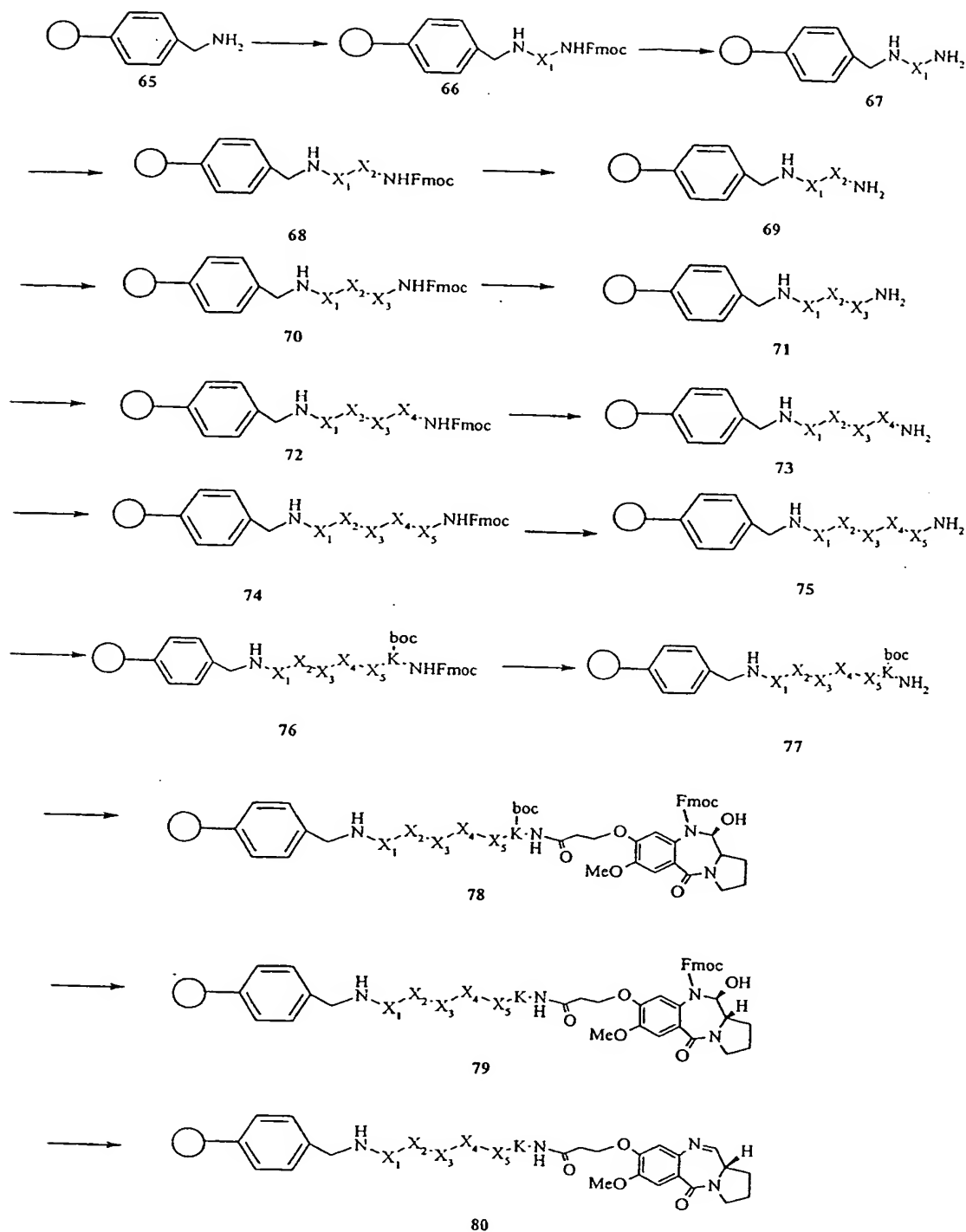


Figure 7

SUBSTITUTE SHEET (RULE 26)

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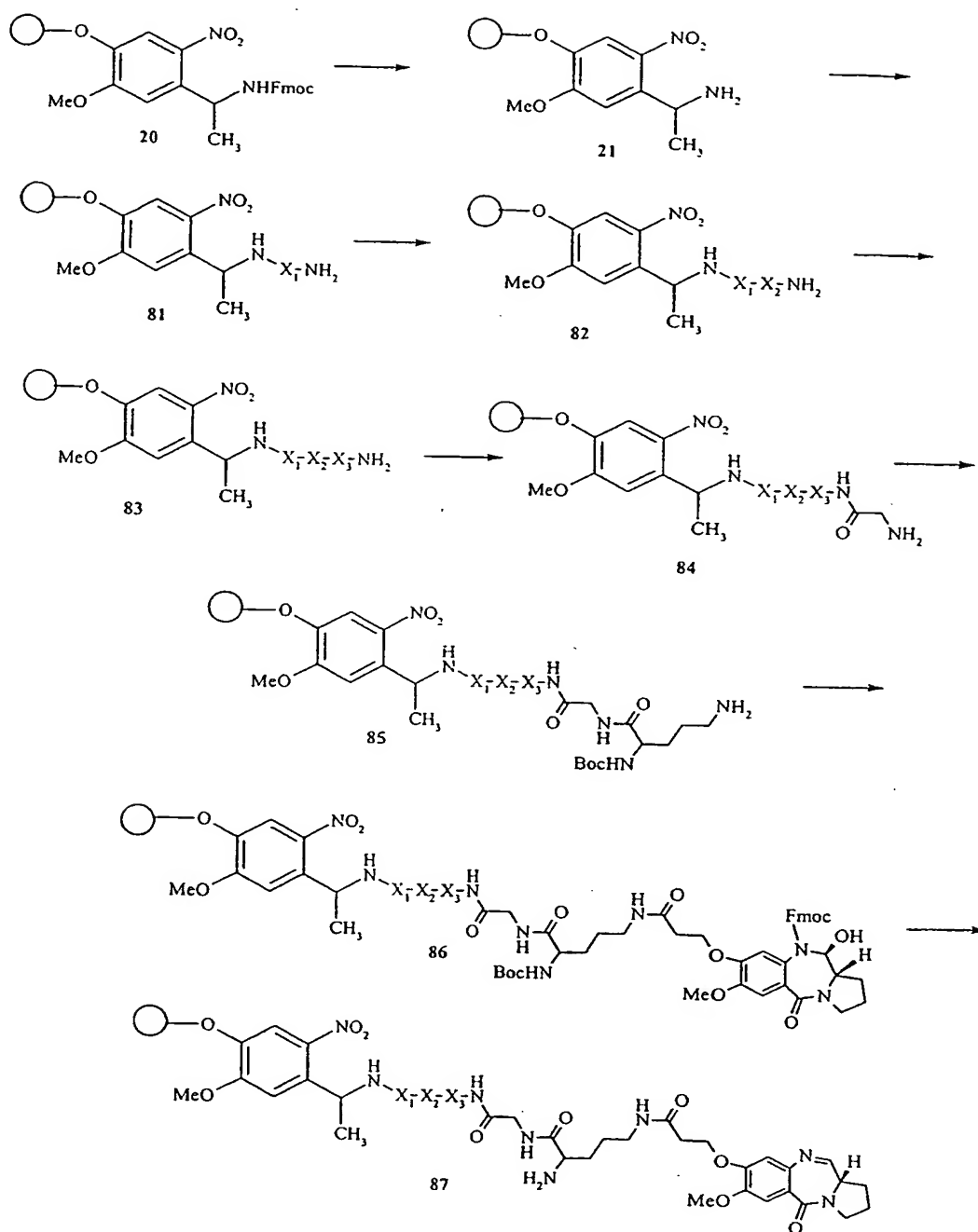


Figure 8

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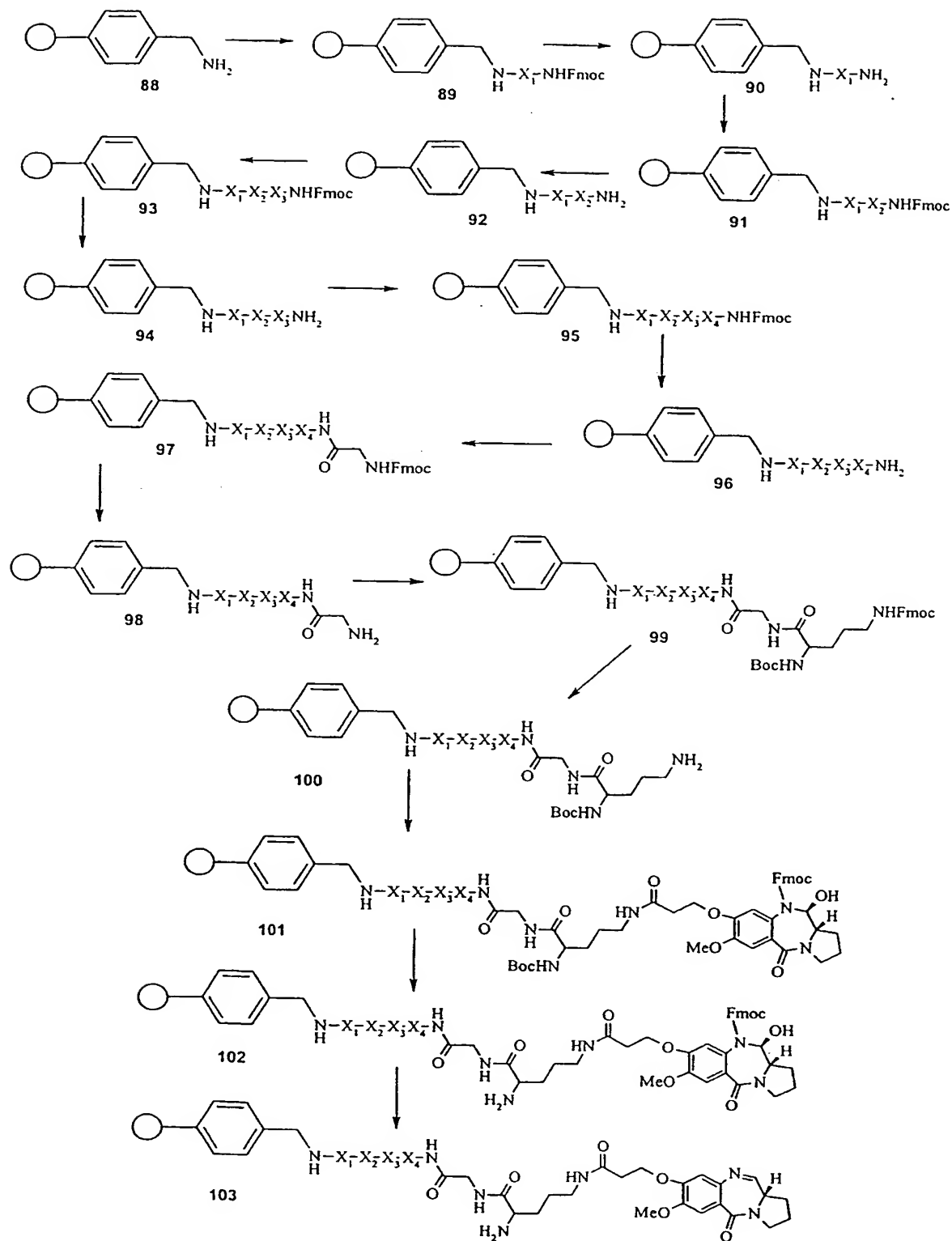


Figure 9

SUBSTITUTE SHEET (RULE 26)

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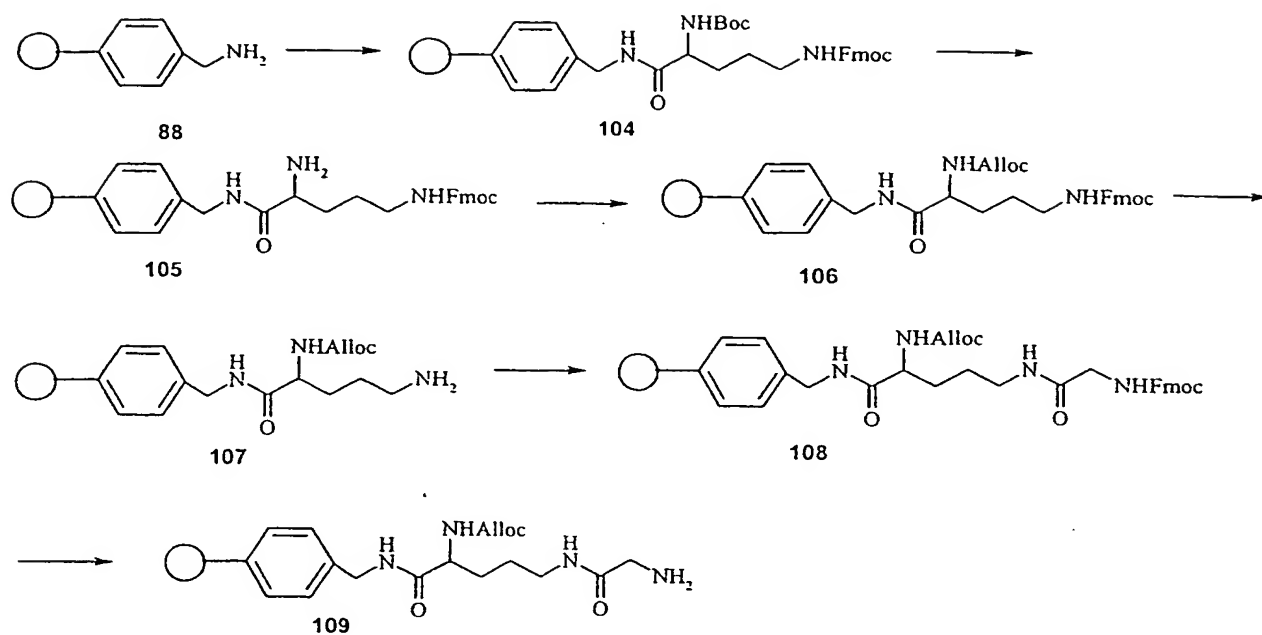


Figure 10a

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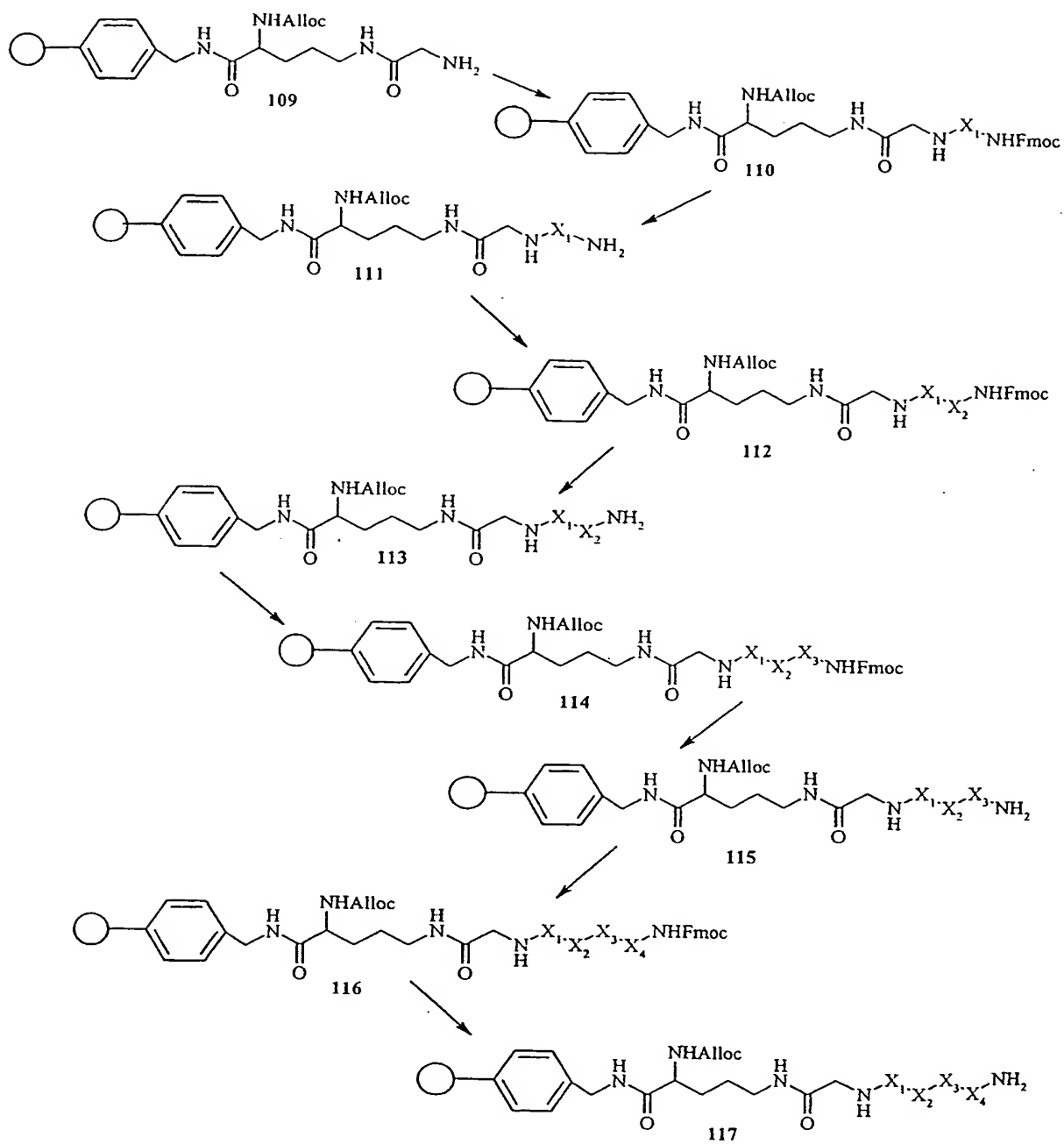


Figure 10b

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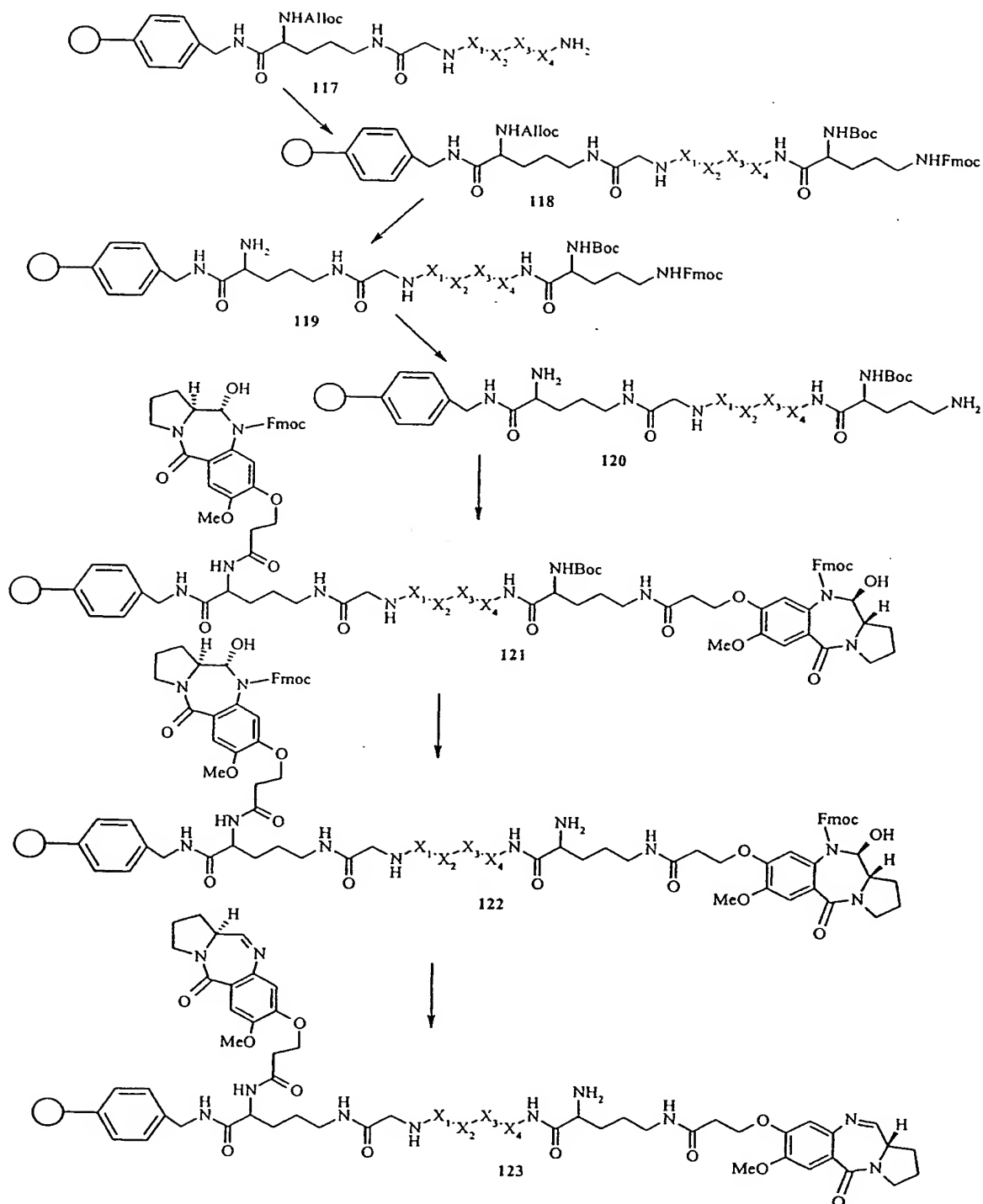


Figure 10c

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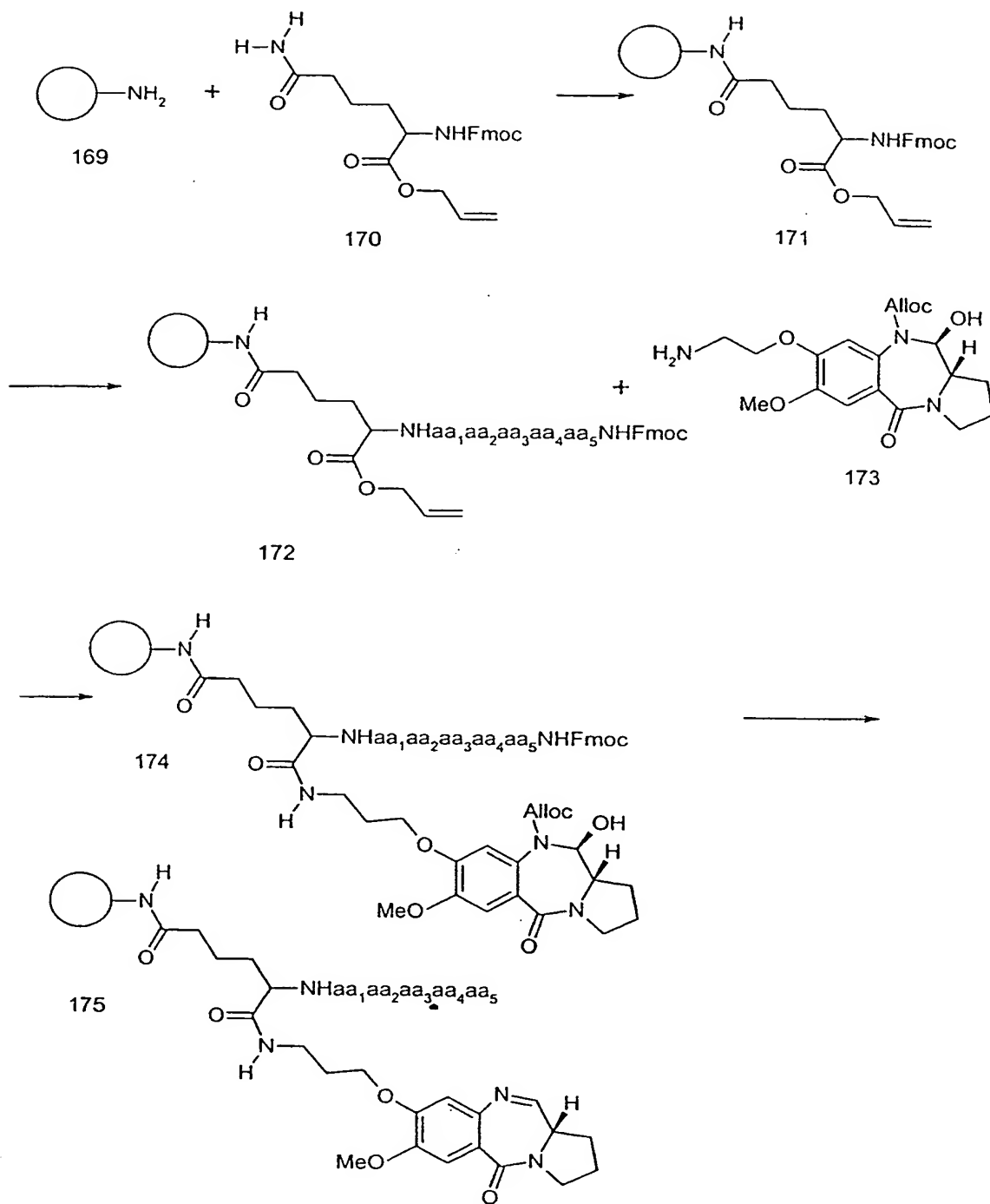


Figure 11

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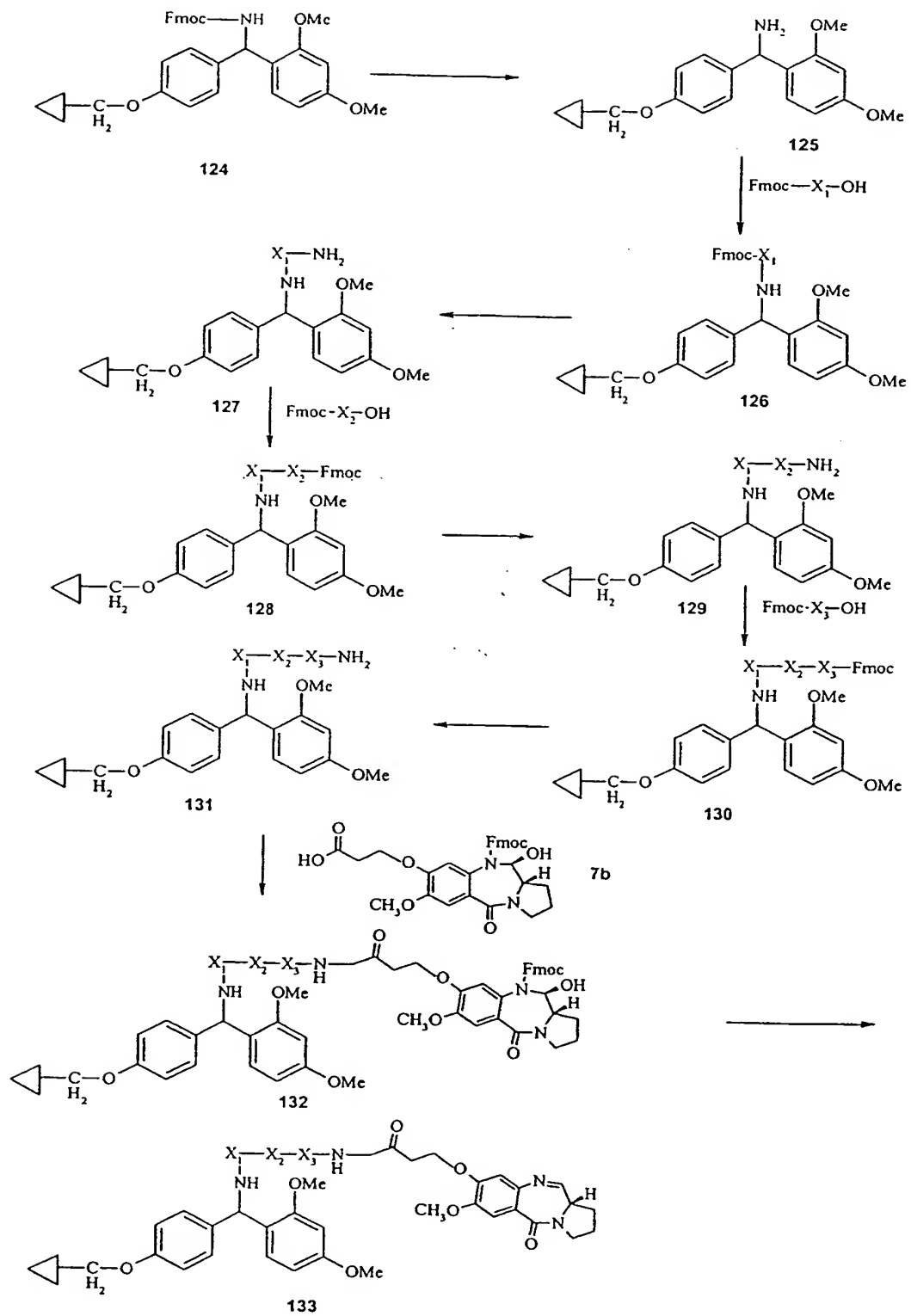
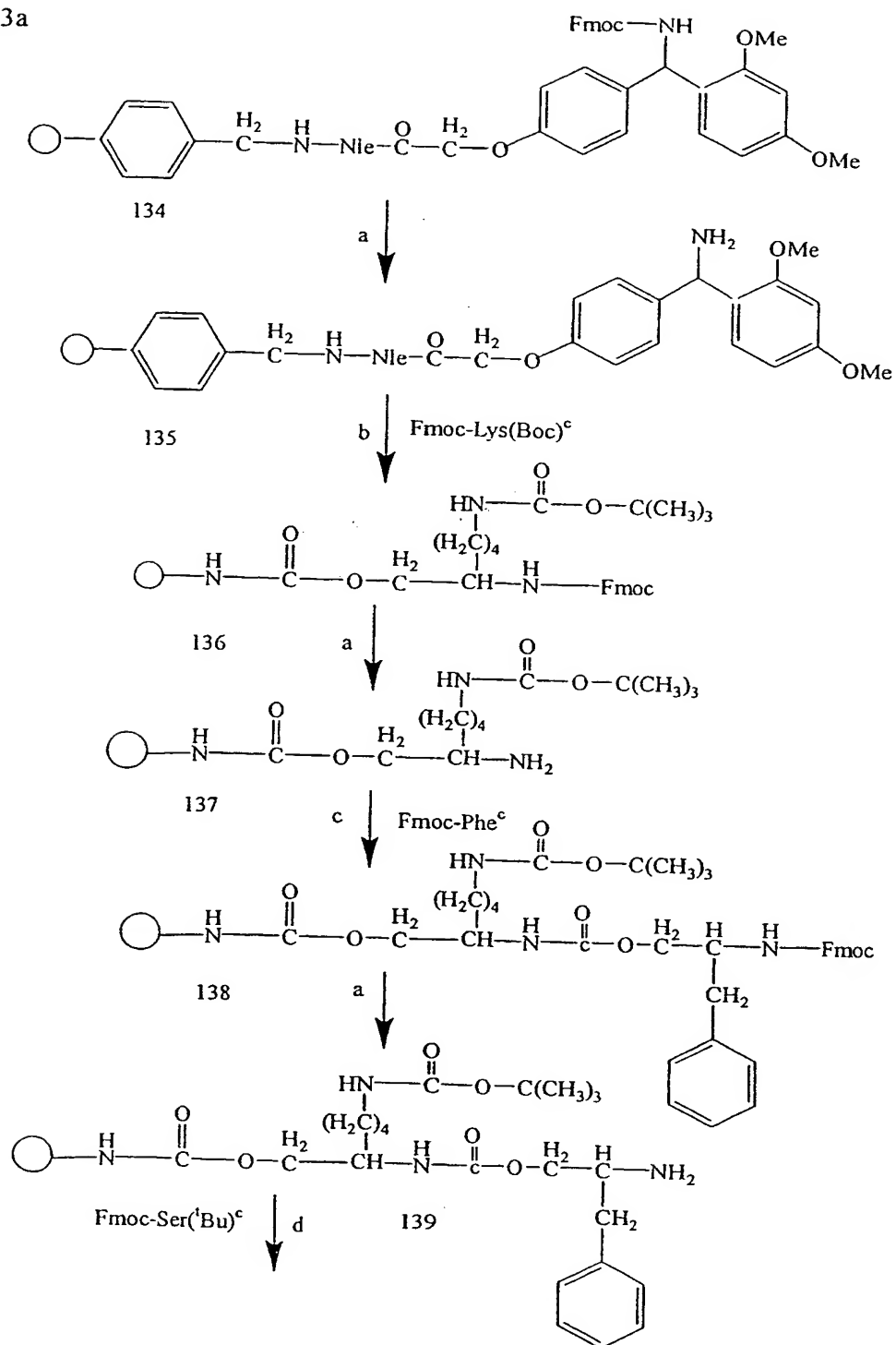


Figure 12

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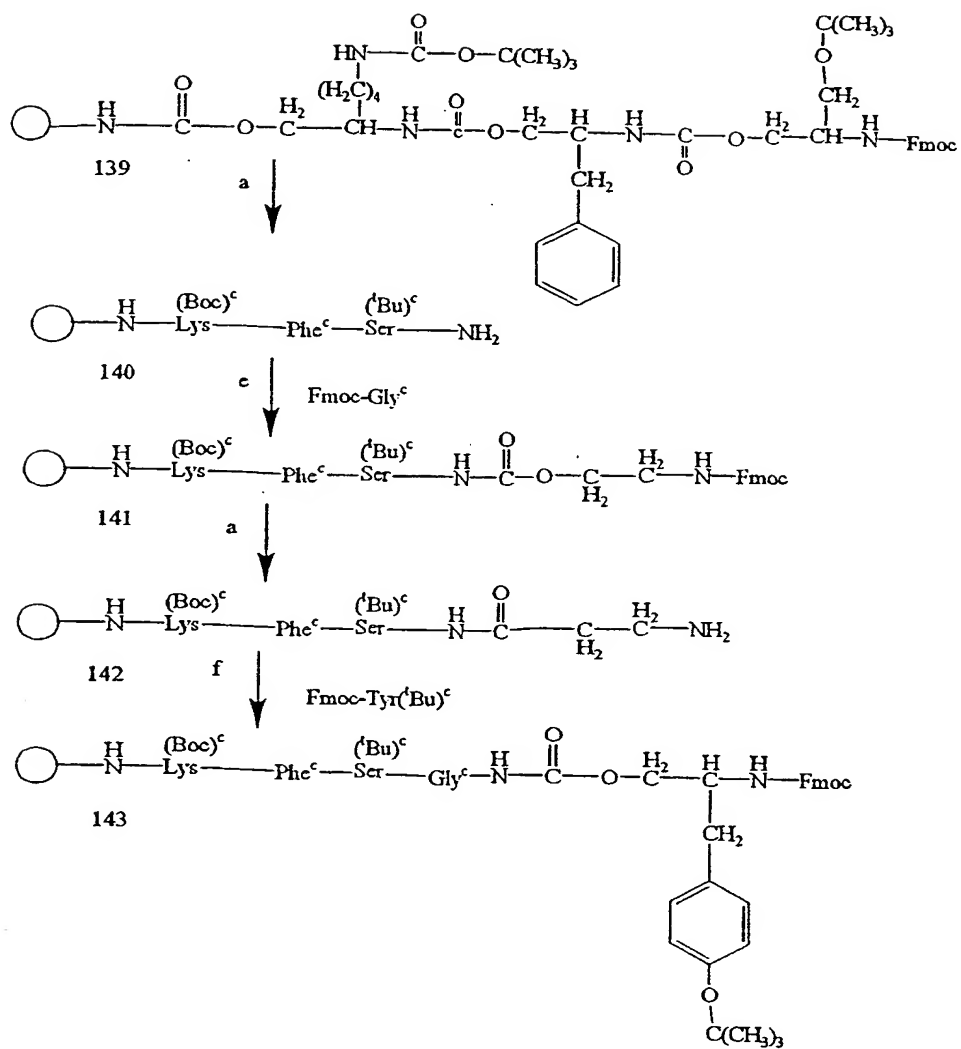
Figure 13a



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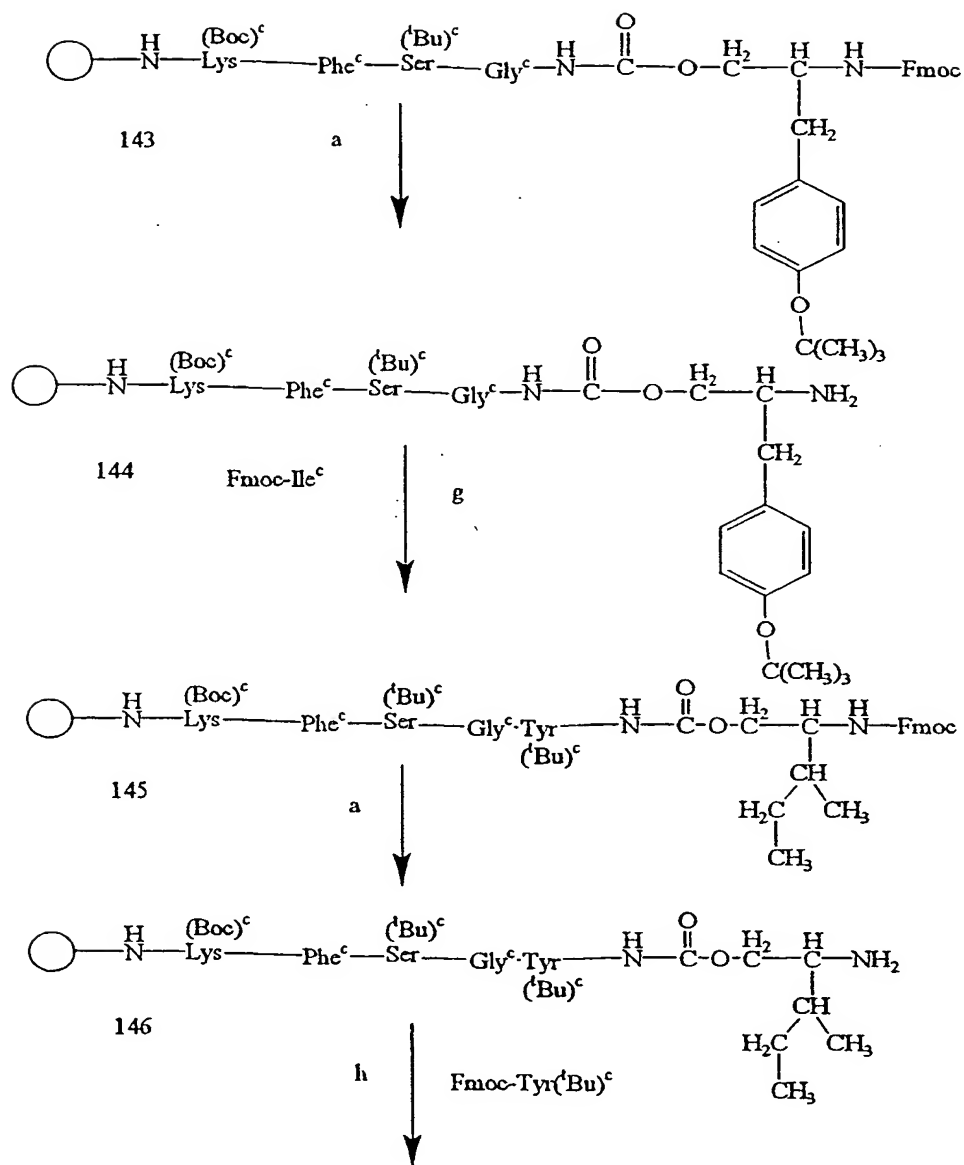
Figure 13b



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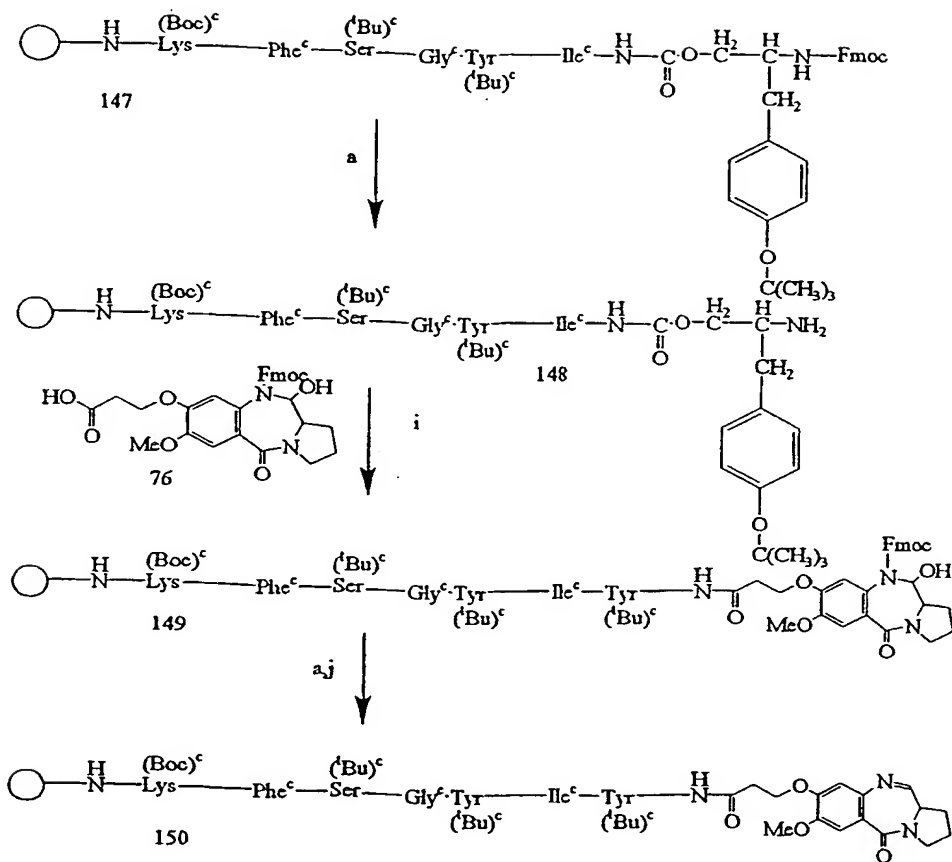
Figure 13c



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Figure 13d



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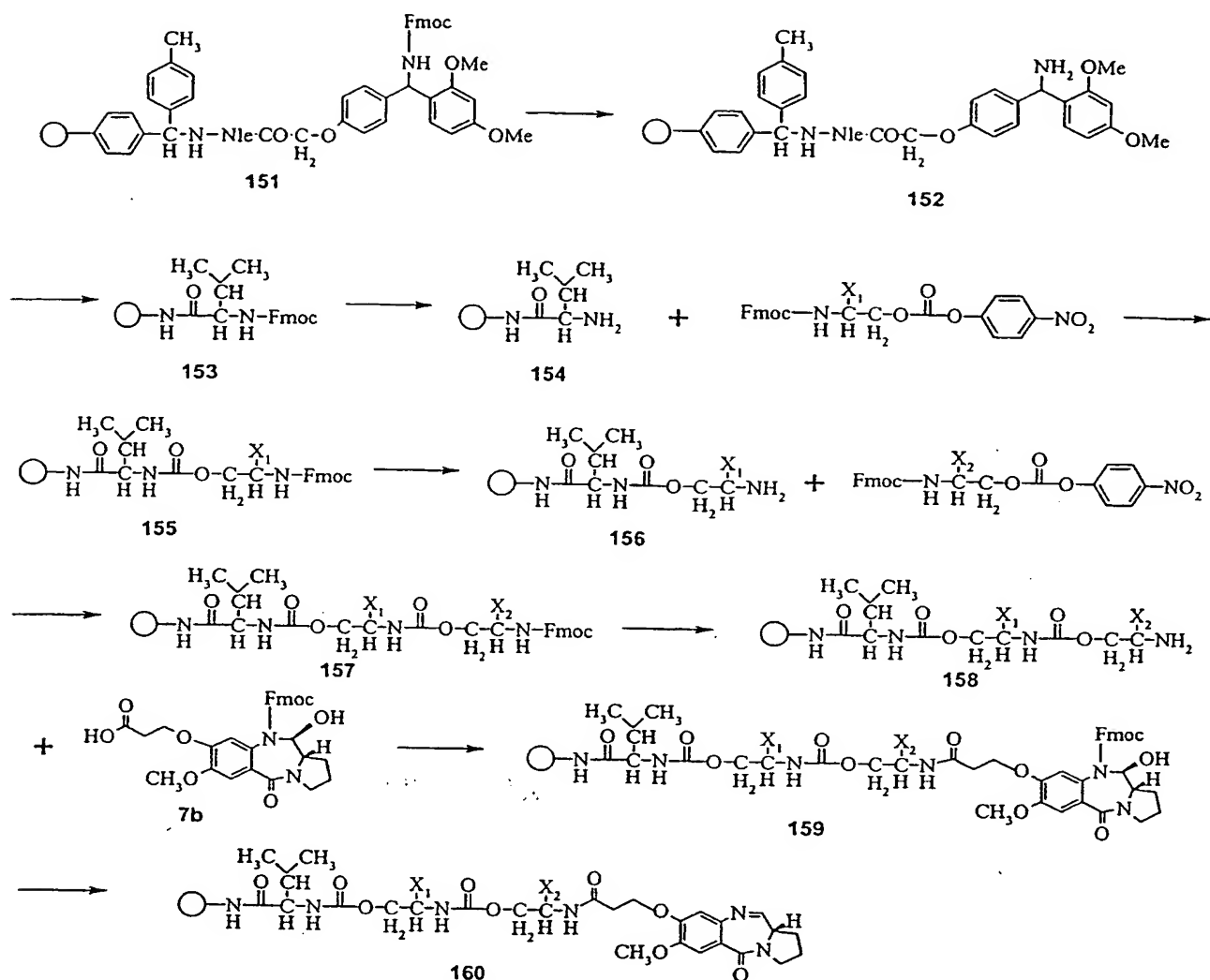


Figure 14

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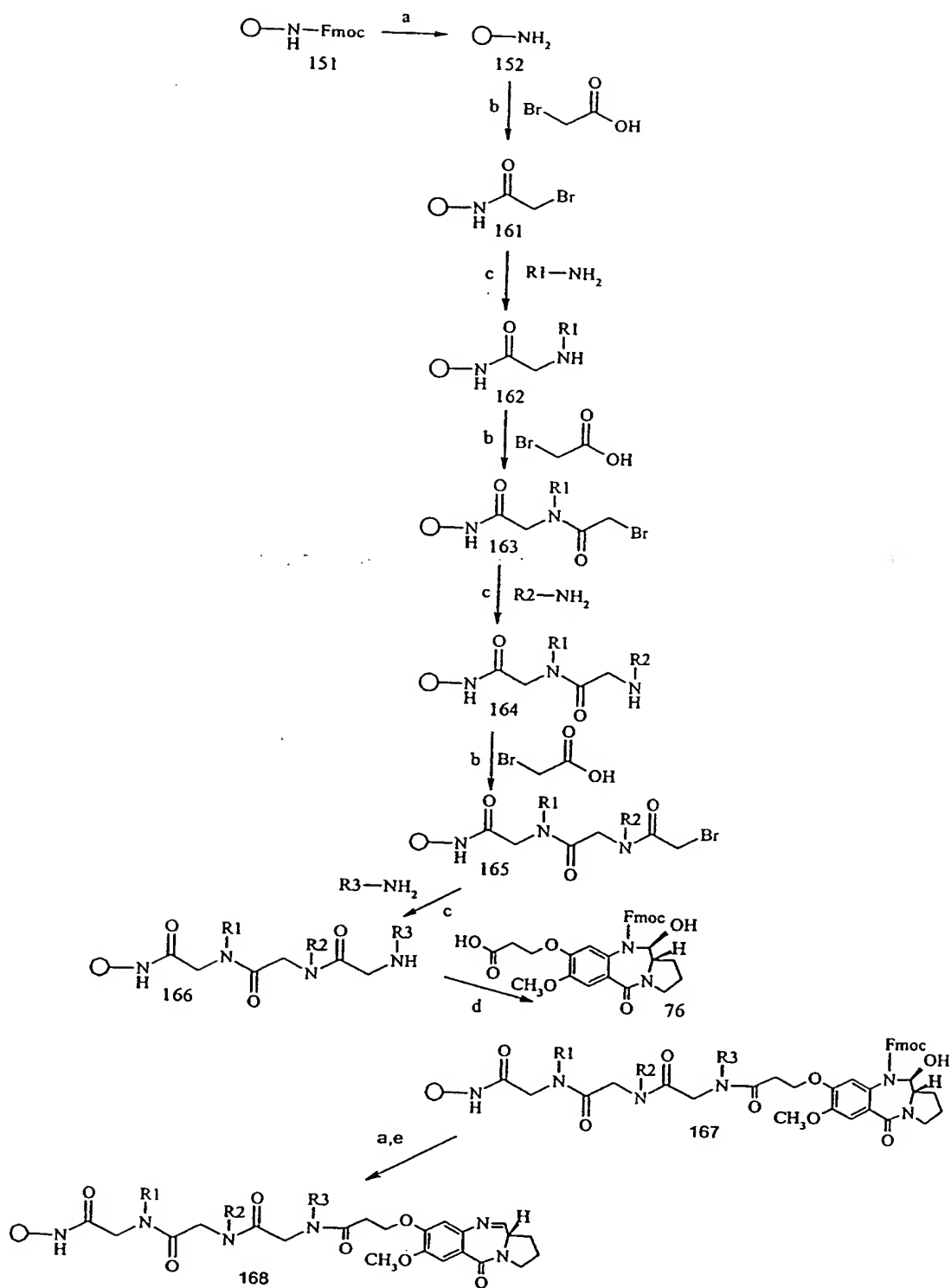


Figure 15

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